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GENES ENCODING LEPIDOPTERAN-ACTIVE TOXINS

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(57) Claim

1. A process for controlling lepidopteran insect pests which comprises contacting said insect pests with an insect-controlling effective amount of *Bacillus thuringiensis* PS81A2, having the identifying characteristics of NRRL B-18457, or *Bacillus thuringiensis* PS81RR1, having the identifying characteristics of NRRL B-18458, or mutants thereof which retain the characteristics of the parent strains.

2. The process, according to claim 1, wherein said mutants are asporogenous mutants and/or phage resistant mutants.

3. The process, according to claim 1, wherein said insect pest is contacted with an insect-controlling effective amount of *Bacillus thuringiensis* PS81A2 or PS81RR1, by incorporating said *Bacillus thuringiensis* PS81A2 or PS81RR1 into a bait granule and placing said granule on or in the soil when planting seed of a plant upon which plant insect pest is known to feed.

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COMPLETE SPECIFICATION

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Complete Specification for the invention entitled:

Novel Bacillus thuringiensis Isolates Active Against
Lepidopteran Pests, and Genes Encoding Lepidopteran-Active
Toxins

The following statement is a full description of this invention, including the
best method of performing it known to me/us

Abstract of the Disclosure

5 Novel Bacillus thuringiensis genes encoding toxins which are active against lepidopteran insects have been cloned from novel lepidopteran-active B. thuringiensis microbes. The DNA encoding the B. thuringiensis toxins can be used to transform various prokaryotic and eukaryotic microbes to express the B. thuringiensis toxins. These recombinant microbes can be used to control lepidopteran insects in various environments.

DESCRIPTION

NOVEL *BACILLUS THURINGIENSIS* ISOLATES
ACTIVE AGAINST LEPIDOPTERAN PESTS
AND GENES ENCODING NOVEL LEPIDOPTERAN-ACTIVE TOXINS

Background of the Invention

The most widely used microbial pesticides are derived from the bacterium Bacillus thuringiensis. This bacterial agent is used to control a wide range of leaf-eating caterpillars and beetles, as well as mosquitos. Bacillus thuringiensis produces a proteinaceous parasporal body or crystal which is toxic upon ingestion by a susceptible insect host. For example, B. thuringiensis subsp. kurstaki HD-1 produces a crystal inclusion consisting of a biotoxin called a delta toxin which is toxic to the larvae of a number of lepidopteran insects. The cloning, sequencing, and expression of this B.t. crystal protein gene in Escherichia coli has been described in the published literature (Schnepf, H.E. and Whitely, H.R. [1981] Proc. Natl. Acad. Sci. USA 78:2893-2897; Schnepf et al.). U.S. Patent 4,448,885 and U.S. Patent 4,467,036 both disclose the expression of B.t. crystal protein in E. coli.

Brief Summary of the Invention

The subject invention concerns novel Bacillus thuringiensis isolates designated B.t. PS81A2 and PS81RR1 which have activity against all lepidopteran pests tested.

Also disclosed and claimed are novel toxin genes which express toxins toxic to lepidopteran insects. These toxin genes can be transferred to suitable hosts via a plasmid vector.

Specifically, the invention comprises novel *B.t.* isolates denoted *B.t.* PS81A2 and PS81RR1, mutants thereof, and novel delta endotoxin genes derived from these *B.t.* isolates which encode proteins which are active against lepidopteran pests. More specifically, the gene in *B.t.* PS81A2 encodes a 133,601 dalton endotoxin, whereas the gene in *B.t.* PS81RR1 encodes a 133,367 dalton endotoxin.

Table 1 discloses the DNA encoding the novel toxin expressed by PS81A2. Table 2 discloses the amino acid sequence of the novel toxin expressed by PS81A2. Table 3 is a composite of Tables 1 and 2. Table 4 discloses the DNA encoding the novel toxin expressed by PS81RR1. Table 5 discloses the amino acid sequence of the novel toxin expressed by PS81RR1. Table 6 is a composite of Tables 4 and 5.

According to a first embodiment of this invention, there is provided a process for controlling lepidopteran insect pests which comprises contacting said insect pests with an insect-controlling effective amount of *Bacillus thuringiensis* PS81A2, having the identifying characteristics of NRRL B-18457, or *Bacillus thuringiensis* PS81RR1, having the identifying characteristics of NRRL B-18458, or mutants thereof which retain the characteristics of the parent strains.

According to a second embodiment of this invention, there is provided a process for controlling soil-inhabiting insect pests of the order Lepidopteran which comprises

- (1) preparing a bait granule comprising *Bacillus thuringiensis* PS81A2 or PS81RR1, or mutants thereof which retain the characteristics of the parent strains, or spores or crystals of *Bacillus thuringiensis* PS81A2 or PS81RR1;

and

- (2) placing said bait granule on or in the soil.

According to a third embodiment of this invention, there is provided a composition of matter comprising *Bacillus thuringiensis* PS81A2 or PS81RR1, or mutants thereof which retain the characteristics of the parent strains, or spores or crystals of *Bacillus thuringiensis* PS81A2 or PS81RR1 in association with an insecticide carrier, wherein said mutants are asporogenous mutants and/or phage resistant mutants.

According to a fourth embodiment of this invention, there is provided a composition of matter comprising *Bacillus thuringiensis* PS81A2 or PS81RR1, or mutants thereof which retain the characteristics of the parent strains, in association with formulation ingredients applied as a seed coating, wherein said mutants are asporogenous mutants and/or phage resistant mutants.

According to a fifth embodiment of this invention, there is provided *Bacillus thuringiensis* PS81A2, having the identifying characteristics of NRRL B-18457, or

mutants thereof which retain the characteristics of the parent strain, having activity against insect pests of the order Lepidoptera.

According to a sixth embodiment of this invention, there is provided *Bacillus thuringiensis* PS81RR1, having the identifying characteristics of NRRL B-18458, or 5 mutants thereof which retain the characteristics of the parent strain, having activity against insect pests of the order Lepidoptera.

According to a seventh embodiment of this invention, there is provided asporogenous and/or phage resistant mutants of *Bacillus thuringiensis* PS81A2 or *Bacillus thuringiensis* PS81RR1 which retain the characteristics of the parent strains.

10 According to an eighth embodiment of this invention, there is provided DNA encoding a *Bacillus thuringiensis* toxin having the amino acid sequences shown in Table 2 or Table 5.

According to a ninth embodiment of this invention, there is provided toxin active against lepidopteran insects having the amino acid sequence shown in Table 2 or Table 5, 15 and mutants thereof which do not alter the protein secondary structure, or if the structure is altered, the biological activity is retained to some degree.

According to a tenth embodiment of this invention, there is provided a recombinant DNA transfer vector comprising DNA having all or part of the nucleotide sequence which codes for the amino acid sequence shown in Table 2 or Table 5.

20 According to an eleventh embodiment of this invention, there is provided a bacterial host transformed to express a *Bacillus thuringiensis* toxin having the amino acid sequence shown in Table 2 or Table 5.

According to a twelfth embodiment of this invention, there is provided a method for controlling lepidopteran insects which comprises administering to said insects or to the 25 environment of said insects a microorganism transformed to express a *Bacillus thuringiensis* toxin having the amino acid sequence shown in Table 2 or Table 5, wherein said microorganism is a species of *Pseudomonas*, *Azotobacter*, *Erwinia*, *Serratia*, *Klebsiella*, *Rhizobium*, *Rhodopseudomonas*, *Methylophilus*, *Agrobacterium*, *Acetobacter*, *Alcaligenes*, *Bacillus*, or *Streptomyces*.

30 According to a thirteenth embodiment of this invention, there is provided an insecticidal composition comprising insecticide containing substantially intact, treated cells having prolonged pesticidal activity when applied to the environment of a target pest, wherein said insecticide is a polypeptide toxic to lepidopteran insects, is intracellular, and is produced as a result of expression of a transformed microbe capable of expressing the 35 *Bacillus thuringiensis* toxin having the amino acid sequence shown in Table 2 or Table 5.

According to a fourteenth embodiment of this invention, there is provided treated, substantially intact unicellular microorganism cells containing an intracellular toxin, which toxin is a result of expression of a *Bacillus thuringiensis* toxin gene toxic to lepidopteran insects which codes for a polypeptide toxin having the amino acid sequence shown in

Table 2 or Table 5, wherein said cells are treated under conditions which prolong the insecticidal activity when said cells are applied to the environment of a target insect.

Brief Description of the Drawings

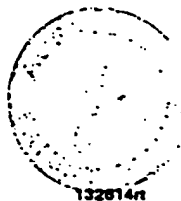
Figure 1 shows agarose gel electrophoresis of plasmid preparations from *B.t.* PS81A2, *B.t.* PS81RR1, and *B.t.* HD-1.

Detailed Disclosure of the Invention

The novel toxin genes of the subject invention were obtained from novel lepidopteran-active *B. thuringiensis* (*B.t.*) isolates designated PS81A2 and PS81RR1.

Characteristics of *B.t.* PS81A2 and PS81RR1

- 10 Colony morphology -- Large colony, dull surface, typical *B.t.*
Vegetative cell morphology -- typical *B.t.*
Flagellar serotype -- 7, aizawai.
Intracellular inclusions -- sporulating cells produce a bipyramidal crystal.



Plasmid preparations -- agarose gel electrophoresis of plasmid preparations distinguishes B.t. PS81A2 and PS81RR1 from B.t. HD-1 and other B.t. isolates. See Figure 1.

Alkali-soluble proteins -- B.t. PS81A2 and PS81RR1 produce 133,601 and 133,367 dalton proteins, respectively.

Unique toxins -- the 133,601 and 133,367 dalton toxins are different from any previously identified.

Activity -- B.t. PS81A2 and PS81RR1 both kill all Lepidoptera tested (Trichoplusia ni, Spodoptera exigua, and Plutella xylostella).

Bioassay procedures:

Spodoptera exigua--dilutions are prepared of a spore and crystal pellet, mixed with USDA Insect Diet (Technical Bulletin 1528, U.S. Department of Agriculture) and poured into small plastic trays. Neonate Spodoptera exigua larvae are placed on the diet mixture and held at 25°C. Mortality is recorded after six days.

Other insects -- dilutions and diet are prepared in the same manner as for the Spodoptera exigua bioassay. Fourth instar larvae are used, and mortality is recorded after eight days.

B. thuringiensis PS81A2, NRRL B-18457, and B. thuringiensis PS81RR1, NRRL B-18458, and mutants thereof, can be cultured using standard known media and fermentation techniques. Upon completion of the fermentation cycle, the bacteria can be harvested by first separating the B.t. spores and crystals from the fermentation broth by means well known in the art. The recovered B.t. spores and crystals can be formulated into a wettable powder, a liquid concentrate, granules or other formulations by the addition of surfactants, dispersants, inert carriers and other components to facilitate handling and

application for particular target pests. The formulation and application procedures are all well known in the art and are used with commercial strains of B. thuringiensis (HD-1) active against Lepidoptera, e.g., caterpillars. B.t. PS81A2 and B.t. PS81RR1, and mutants thereof, can be used to control lepidopteran pests.

A subculture of B.t. PS81A2 and PS81RR1 and the E. coli hosts harboring the toxin genes of the invention, were deposited in the permanent collection of the Northern Research Laboratory, U.S. Department of Agriculture, Peoria, Illinois, USA. The accession numbers and deposit dates are as follows:

<u>Subculture</u>	<u>Accession Number</u>	<u>Deposit Date</u>
<u>B.t.</u> PS81A2	NRRL B-18457	March ⁷ 14 , 1989
<u>B.t.</u> PS81RR1	NRRL B-18458	March ⁷ 14 , 1989
<u>E. coli</u> (NM522)(pMYC389)	NRRL B-18448	February 24, 1989
<u>E. coli</u> (NM522)(pMYC390)	NRRL B-18449	February 24, 1989

The subject cultures have been deposited under conditions that assure that access to the cultures will be available during the pendency of this patent application to one determined by the Commissioner of Patents and Trademarks to be entitled thereto under 37 CFR 1.14 and 35 USC 122. The deposits are available as required by foreign patent laws in countries wherein counterparts of the subject application, or its progeny, are filed. However, it should be understood that the availability of a deposit does not constitute a license to practice the subject invention in derogation of patent rights granted by governmental action.

Further, the subject culture deposits will be stored and made available to the public in accord with the provisions of the Budapest Treaty for the Deposit of Microorganisms, i.e., they will be stored with all the care necessary to keep



them viable and uncontaminated for a period of at least five years after the most recent request for the furnishing of a sample of the deposit, and in any case, for a period of at least 30 (thirty) years after the date of deposit or for the enforceable life of any patent which may issue disclosing the cultures. The depositor acknowledges the duty to replace the deposits should the depository be unable to furnish a sample when requested, due to the condition of the deposit(s). All restrictions on the availability to the public of the subject culture deposits will be irrevocably removed upon the granting of a patent disclosing them.

The toxin genes of the subject invention can be introduced into a wide variety of microbial hosts. Expression of the toxin gene results, directly or indirectly, in the intracellular production and maintenance of the pesticide. With suitable hosts, e.g., Pseudomonas, the microbes can be applied to the situs of lepidopteran insects where they will proliferate and be ingested by the insects. The result is a control of the unwanted insects. Alternatively, the microbe hosting the toxin gene can be treated under conditions that prolong the activity of the toxin produced in the cell. The treated cell then can be applied to the environment of target pest(s). The resulting product retains the toxicity of the B.t. toxin.

Where the B.t. toxin gene is introduced via a suitable vector into a microbial host, and said host is applied to the environment in a living state, it is essential that certain host microbes be used. Microorganism hosts are selected which are known to occupy the "phytosphere" (phylloplane, phyllosphere, rhizosphere, and/or rhizoplane) of one or more crops of interest. These microorganisms are selected so as to be capable of successfully competing in the particular environment (crop and other insect habitats) with the wild-type microorganisms, provide for stable maintenance and expression of the gene

expressing the polypeptide pesticide, and, desirably, provide for improved protection of the pesticide from environmental degradation and inactivation.

A large number of microorganisms are known to inhabit the phylloplane (the surface of the plant leaves) and/or the rhizosphere (the soil surrounding plant roots) of a wide variety of important crops. These microorganisms include bacteria, algae, and fungi. Of particular interest are microorganisms, such as bacteria, e.g., genera Bacillus, Pseudomonas, Erwinia, Serratia, Klebsiella, Xanthomonas, Streptomyces, Rhizobium, Rhodopseudomonas, Methylophilus, Agrobacterium, Acetobacter, Lactobacillus, Arthrobacter, Azotobacter, Leuconostoc, and Alcaligenes; fungi, particularly yeast, e.g., genera Saccharomyces, Cryptococcus, Kluyveromyces, Sporobolomyces, Rhodotorula, and Aureobasidium. Of particular interest are such phytosphere bacterial species as Pseudomonas syringae, Pseudomonas fluorescens, Serratia marcescens, Acetobacter xylinum, Agrobacterium tumefaciens, Rhodopseudomonas spheroides, Xanthomonas campestris, Rhizobium melioli, Alcaligenes entrophus, and Azotobacter vinlandii; and phytosphere yeast species such as Rhodotorula rubra, R. glutinis, R. marina, R. aurantiaca, Cryptococcus albidus, C. diffluens, C. laurentii, Saccharomyces rosei, S. pretoriensis, S. cerevisiae, Sporobolomyces roseus, S. odor, Kluyveromyces veronae, and Aureobasidium pollulans. Of particular interest are the pigmented microorganisms.

A wide variety of ways are available for introducing a B.t. gene expressing a toxin into the microorganism host under conditions which allow for stable maintenance and expression of the gene. One can provide for DNA constructs which include the transcriptional and translational regulatory signals for expression of the toxin gene, the toxin gene under their regulatory control and a DNA sequence homologous with a sequence in the host organism, whereby integration will occur, and/or a replication system which is functional in the host, whereby integration or stable maintenance will occur.

The transcriptional initiation signals will include a promoter and a transcriptional initiation start site. In some instances, it may be desirable to provide for regulative expression of the toxin, where expression of the toxin will only occur after release into the environment. This can be achieved with operators or a region binding to an activator or enhancers, which are capable of induction upon a change in the physical or chemical environment of the microorganisms. For example, a temperature sensitive regulatory region may be employed, where the organisms may be grown up in the laboratory without expression of a toxin, but upon release into the environment, expression would begin. Other techniques may employ a specific nutrient medium in the laboratory, which inhibits the expression of the toxin, where the nutrient medium in the environment would allow for expression of the toxin. For translational initiation, a ribosomal binding site and an initiation codon will be present.

Various manipulations may be employed for enhancing the expression of the messenger RNA, particularly by using an active promoter, as well as by employing sequences, which enhance the stability of the messenger RNA. The transcriptional and translational termination region will involve stop codon(s), a terminator region, and optionally, a polyadenylation signal. A hydrophobic "leader" sequence may be employed at the amino terminus of the translated polypeptide sequence in order to promote secretion of the protein across the inner membrane.

In the direction of transcription, namely in the 5' to 3' direction of the coding or sense sequence, the construct will involve the transcriptional regulatory region, if any, and the promoter, where the regulatory region may be either 5' or 3' of the promoter, the ribosomal binding site, the initiation codon, the structural gene having an open reading frame in phase with the initiation codon, the stop codon(s), the polyadenylation signal sequence, if any, and the terminator region. This sequence as a double strand may be used by itself for

transformation of a microorganism host, but will usually be included with a DNA sequence involving a marker, where the second DNA sequence may be joined to the toxin expression construct during introduction of the DNA into the host.

By a marker is intended a structural gene which provides for selection of those hosts which have been modified or transformed. The marker will normally provide for selective advantage, for example, providing for biocide resistance, e.g., resistance to antibiotics or heavy metals; complementation, so as to provide prototrophy to an auxotrophic host, or the like. Preferably, complementation is employed, so that the modified host may not only be selected, but may also be competitive in the field. One or more markers may be employed in the development of the constructs, as well as for modifying the host. The organisms may be further modified by providing for a competitive advantage against other wild-type microorganisms in the field. For example, genes expressing metal chelating agents, e.g., siderophores, may be introduced into the host along with the structural gene expressing the toxin. In this manner, the enhanced expression of a siderophore may provide for a competitive advantage for the toxin-producing host, so that it may effectively compete with the wild-type microorganisms and stably occupy a niche in the environment.

Where no functional replication system is present, the construct will also include a sequence of at least 50 basepairs (bp), preferably at least about 100 bp, and usually not more than about 1000 bp of a sequence homologous with a sequence in the host. In this way, the probability of legitimate recombination is enhanced, so that the gene will be integrated into the host and stably maintained by the host. Desirably, the toxin gene will be in close proximity to the gene providing for complementation as well as the gene providing for the competitive advantage. Therefore, in the event that a toxin gene is lost, the resulting organism will be likely to also lose the complementing gene and/or the gene

providing for the competitive advantage, so that it will be unable to compete in the environment with the gene retaining the intact construct.

5 A large number of transcriptional regulatory regions are available from a wide variety of microorganism hosts, such as bacteria, bacteriophage, cyanobacteria, algae, fungi, and the like. Various transcriptional regulatory regions include the regions associated with the trp gene, lac gene, gal gene, the lambda left and right promoters, the Tac promoter, the naturally-occurring promoters associated with the toxin gene, where functional in the host. See for example, U.S. Patent Nos. 4,332,898, 4,342,832 and 4,356,270. The termination
10 region may be the termination region normally associated with the transcriptional initiation region or a different transcriptional initiation region, so long as the two regions are compatible and functional in the host.

Where stable episomal maintenance or integration is desired, a plasmid will be employed which has a replication system which is functional in the host.
15 The replication system may be derived from the chromosome, an episomal element normally present in the host or a different host, or a replication system from a virus which is stable in the host. A large number of plasmids are available, such as pBR322, pACYC184, RSF1010, pRO1614, and the like. See for example, Olson et al., (1982) J. Bacteriol. 150:6069, and Bagdasarian et al.,
20 (1981) Gene 16:237, and U.S. Patent Nos. 4,356,270, 4,362,817, and 4,371,625.

The B.t. gene can be introduced between the transcriptional and translational initiation region and the transcriptional and translational termination region, so as to be under the regulatory control of the initiation region. This construct will be included in a plasmid, which will include at least one replication
25 system, but may include more than one, where one replication system is employed for cloning during the development of the plasmid and the second replication system is necessary for functioning in the ultimate host. In addition, one or more markers may be present, which have been described previously.

Where integration is desired, the plasmid will desirably include a sequence homologous with the host genome.

The transformants can be isolated in accordance with conventional ways, usually employing a selection technique, which allows for selection of the desired organism as against unmodified organisms or transferring organisms, when present. The transformants then can be tested for pesticidal activity.

Suitable host cells, where the pesticide-containing cells will be treated to prolong the activity of the toxin in the cell when the then treated cell is applied to the environment of target pest(s), may include either prokaryotes or eukaryotes, normally being limited to those cells which do not produce substances toxic to higher organisms, such as mammals. However, organisms which produce substances toxic to higher organisms could be used, where the toxin is unstable or the level of application sufficiently low as to avoid any possibility of toxicity to a mammalian host. As hosts, of particular interest will be the prokaryotes and the lower eukaryotes, such as fungi. Illustrative prokaryotes, both Gram-negative and -positive, include Enterobacteriaceae, such as Escherichia, Erwinia, Shigella, Salmonella, and Proteus; Bacillaceae; Rhizobiceae, such as Rhizobium; Spirillaceae, such as photobacterium, Zymomonas, Serratia, Aeromonas, Vibrio, Desulfovibrio, Spirillum; Lactobacillaceae; Pseudomonadaceae, such as Pseudomonas and Acetobacter; Azotobacteraceae, Actinomycetales, and Nitrobacteraceae. Among eukaryotes are fungi, such as Phycomycetes and Ascomycetes, which includes yeast, such as Saccharomyces and Schizosaccharomyces; and Basidiomycetes yeast, such as Rhodotorula, Aureobasidium, Sporobolomyces, and the like.

Characteristics of particular interest in selecting a host cell for purposes of production include ease of introducing the B.t. gene into the host, availability of expression systems, efficiency of expression, stability of the pesticide in the host, and the presence of auxiliary genetic capabilities. Characteristics of interest

for use as a pesticide microcapsule include protective qualities for the pesticide, such as thick cell walls, pigmentation, and intracellular packaging or formation of inclusion bodies; leaf affinity; lack of mammalian toxicity; attractiveness to pests for ingestion; ease of killing and fixing without damage to the toxin; and the like. Other considerations include ease of formulation and handling, economics, storage stability, and the like.

Host organisms of particular interest include yeast, such as Rhodotorula sp., Aureobasidium sp., Saccharomyces sp., and Sporobolomyces sp.; phylloplane organisms such as Pseudomonas sp., Erwinia sp. and Flavobacterium sp.; or such other organisms as Escherichia, Lactobacillus sp., Bacillus sp., Streptomyces sp., and the like. Specific organisms include Pseudomonas aeruginosa, Pseudomonas fluorescens, Saccharomyces cerevisiae, Bacillus thuringiensis, Escherichia coli, Bacillus subtilis, Streptomyces lividans and the like.

The cell will usually be intact and be substantially in the proliferative form when treated, rather than in a spore form, although in some instances spores may be employed.

Treatment of the microbial cell, e.g., a microbe containing the B.t. toxin gene, can be by chemical or physical means, or by a combination of chemical and/or physical means, so long as the technique does not deleteriously affect the properties of the toxin, nor diminish the cellular capability in protecting the toxin. Examples of chemical reagents are halogenating agents, particularly halogens of atomic no. 17-80. More particularly, iodine can be used under mild conditions and for sufficient time to achieve the desired results. Other suitable techniques include treatment with aldehydes, such as formaldehyde and glutaraldehyde; anti-infectives, such as zephiran chloride and cetylpyridinium chloride; alcohols, such as isopropyl and ethanol; various histologic fixatives, such as Lugol iodine, Bouin's fixative, and Helly's fixative (See: Humason, Gretchen L., Animal Tissue Techniques, W.H. Freeman and Company, 1967); or a combination of physical

(heat) and chemical agents that preserve and prolong the activity of the toxin produced in the cell when the cell is administered to the host animal. Examples of physical means are short wavelength radiation such as gamma-radiation and X-radiation, freezing, UV irradiation, lyophilization, and the like.

5 The cells generally will have enhanced structural stability which will enhance resistance to environmental conditions. Where the pesticide is in a proform, the method of inactivation should be selected so as not to inhibit processing of the proform to the mature form of the pesticide by the target pest pathogen. For example, formaldehyde will crosslink proteins and could inhibit
10 processing of the proform of a polypeptide pesticide. The method of inactivation or killing retains at least a substantial portion of the bio-availability or bioactivity of the toxin.

15 The cellular host containing the B.t. insecticidal gene may be grown in any convenient nutrient medium, where the DNA construct provides a selective advantage, providing for a selective medium so that substantially all or all of the cells retain the B.t. gene. These cells may then be harvested in accordance with conventional ways. Alternatively, the cells can be treated prior to harvesting.

20 The B.t. cells may be formulated in a variety of ways. They may be employed as wettable powders, granules or dusts, by mixing with various inert materials, such as inorganic minerals (phyllosilicates, carbonates, sulfates, phosphates, and the like) or botanical materials (powdered corncobs, rice hulls, walnut shells, and the like). The formulations may include spreader-sticker adjuvants, stabilizing agents, other pesticidal additives, or surfactants. Liquid
25 formulations may be aqueous-based or non-aqueous and employed as foams, gels, suspensions, emulsifiable concentrates, or the like. The ingredients may include rheological agents, surfactants, emulsifiers, dispersants, or polymers.

 The pesticidal concentration will vary widely depending upon the nature of the particular formulation, particularly whether it is a concentrate or to be

used directly. The pesticide will be present in at least 1% by weight and may be 100% by weight. The dry formulations will have from about 1-95% by weight of the pesticide while the liquid formulations will generally be from about 1-60% by weight of the solids in the liquid phase. The formulations will generally have from about 10^2 to about 10^4 cells/mg. These formulations will be administered at about 50 mg (liquid or dry) to 1 kg or more per hectare.

The formulations can be applied to the environment of the lepidopteran pest(s), e.g., plants, soil or water, by spraying, dusting, sprinkling, or the like.

Mutants of PS81A2 and PS81RR1 can be made by procedures well known in the art. For example, an asporogenous mutant can be obtained through ethylmethane sulfonate (EMS) mutagenesis of PS81A2 and PS81RR1. The mutants can be made using ultraviolet light and nitrosoguanidine by procedures well known in the art.

A smaller percentage of the asporogenous mutants will remain intact and not lyse for extended fermentation periods; these strains are designated lysis minus (-). Lysis minus strains can be identified by screening asporogenous mutants in shake flask media and selecting those mutants that are still intact and contain toxin crystals at the end of the fermentation. Lysis minus strains are suitable for a cell fixation process that will yield a protected, encapsulated toxin protein.

To prepare a phage resistant variant of said asporogenous mutant, an aliquot of the phage lysate is spread onto nutrient agar and allowed to dry. An aliquot of the phage sensitive bacterial strain is then plated directly over the dried lysate and allowed to dry. The plates are incubated at 30°C. The plates are incubated for 2 days and, at that time, numerous colonies could be seen growing on the agar. Some of these colonies are picked and subcultured onto nutrient agar plates. These apparent resistant cultures are tested for resistance by cross streaking with the phage lysate. A line of the phage lysate is streaked

on the plate and allowed to dry. The presumptive resistant cultures are then streaked across the phage line. Resistant bacterial cultures show no lysis anywhere in the streak across the phage line after overnight incubation at 30°C. The resistance to phage is then reconfirmed by plating a lawn of the resistant culture onto a nutrient agar plate. The sensitive strain is also plated in the same manner to serve as the positive control. After drying, a drop of the phage lysate is plated in the center of the plate and allowed to dry. Resistant cultures showed no lysis in the area where the phage lysate has been placed after incubation at 30°C for 24 hours.

Following are examples which illustrate procedures, including the best mode, for practicing the invention. These examples should not be construed as limiting. All percentages are by weight and all solvent mixture proportions are by volume unless otherwise noted.

Example 1 - Culturing *B.t.* PS81A2 and PS81RR1

A subculture of *B.t.* PS81A2 and PS81RR1, or mutants thereof, can be used to inoculate the following medium, a peptone, glucose, salts medium.

Bacto Peptone	7.5 g/l
Glucose	1.0 g/l
KH ₂ PO ₄	3.4 g/l
K ₂ HPO ₄	4.35 g/l
Salt Solution	5.0 ml/l
CaCl ₂ Solution	5.0 ml/l

Salts Solution (100 ml)

MgSO₄·7H₂O 2.46 g

MnSO₄·H₂O 0.04 g

ZnSO₄·7H₂O 0.28 g

5 FeSO₄·7H₂O 0.40 g

CaCl₂ Solution (100 ml)

CaCl₂·2H₂O 3.66 g

pH 7.2

10 The salts solution and CaCl₂ solution are filter-sterilized and added to the autoclaved and cooked broth at the time of inoculation. Flasks are incubated at 30°C on a rotary shaker at 200 rpm for 64 hr.

15 The above procedure can be readily scaled up to large fermentors by procedures well known in the art.

The B.t. spores and/or crystals, obtained in the above fermentation, can be isolated by procedures well known in the art. A frequently-used procedure is to subject the harvested fermentation broth to separation techniques, e.g., centrifugation.

20 Example 2 - Cloning of Novel Toxin Gene From Isolate PS81A2 and Transformation into *Escherichia coli*

25 Total cellular DNA was prepared from B.t. cells grown to a low optical density (OD₆₀₀ = 1.0). The cells were recovered by centrifugation and protoplasted in TES buffer (30 mM Tris-Cl, 10 mM ethylenediaminetetraacetic acid [EDTA], 50 mM NaCl, pH = 8.0) containing 20% sucrose and 50 mg/ml lysozyme. The protoplasts were lysed by addition of sodium dodecyl sulfate (SDS) to a final concentration of 4%. The cellular material was precipitated

overnight at 4°C in 100 mM (final concentration) neutral potassium chloride. The supernate was extracted twice with phenol/chloroform (1:1). The DNA was precipitated with ethanol and purified by isopycnic banding on a cesium gradient.

5 Total cellular DNA from PS81A2 and B.t.k. HD-1 was digested with EcoRI and separated by electrophoresis on a 0.8% Agarose-TAE-buffered gel. A Southern blot of the gel was probed with the NsiI to NsiI fragment of the toxin gene contained in plasmid pM3,130-7 of NRRL B-18332 and the NsiI to KpnI fragment of the "4.5 Kb class" toxin gene (Kronstad and Whitely [1986] Gene USA 43:29-40). These two fragments were combined and used as the probe. Results show that hybridizing fragments of PS81A2 are distinct from those of HD-1. Specifically, a 3.0 Kb hybridizing band in PS81A2 was detected instead of the 3.8 Kb and 1.8 Kb hybridizing bands seen in HD-1.

15 Two hundred micrograms of PS81A2 total cellular DNA was digested with EcoRI and separated by electrophoresis on a preparative 0.8% Agarose-TAE gel. The 2.5 Kb to 3.5 Kb region of the gel was cut out and the DNA from it was electroeluted and concentrated using an ELUTIP™-d (Schleicher and Schuell, Keene, NH) ion exchange column. The isolated EcoRI fragments were ligated to LAMBDA ZAP™ EcoRI arms (Stratagene Cloning Systems, La Jolla, CA) and packaged using Gigapak GOLD™ (Stratagene) extracts. The packaged recombinant phage were plated with E. coli strain BB4 (Stratagene) to give high plaque density. The plaques were screened by standard nucleic acid hybridization procedure with radiolabeled probe. The plaques that hybridized were purified and re-screened at a lower plaque density. The resulting purified phage were grown with R408 M13 helper phage (Stratagene) and the recombinant BlueScript™ (Stratagene) plasmid was automatically excised and packaged. The "phagemid" was re-infected in XL1-Blue E. coli cells (Stratagene) as part of the automatic excision process. The infected XL1-Blue cells were screened for ampicillin resistance and the resulting colonies were analyzed by standard

20

25

miniprep procedure to find the desired plasmid. The plasmid, designated pM6,31-1, contains an approximate 3.0 Kb EcoRI insert and was sequenced using Stratagene's T7 and T3 primers plus a set of existing B.t. endotoxin gene oligonucleotide primers. About 1.8 Kb of the toxin gene was sequenced, and data analysis comparing PS81A2 to other cloned B.t. endotoxin genes showed that the PS81A2 sequence was unique. A synthetic oligonucleotide (CAGATCCACGAGGCTTATCTTCCAGAACTAC) was constructed to one of the regions in the PS81A2 sequence that was least homologous relative to other exiting B.t. endotoxin genes.

PS81A2 total cellular DNA partially digested with Sau3A and fractionated by electrophoresis into a mixture of 9-23 Kb fragments on a 0.6% agarose-TAE gel was ligated into Lambda DASHTM (Stratagene). The packaged phage at a high titer were plated on P2392 E. coli cells (Stratagene) and screened using the radiolabeled synthetic oligonucleotide (aforementioned) as a nucleic acid hybridization probe. Hybridizing plaques were rescreened at a lower plaque density. A single purified hybridizing plaque was used to infect P2392 E. coli cells in liquid culture for preparation of phage for DNA isolation. DNA was isolated by standard procedures. Preparative amounts of recombinant phage DNA were digested with SalI (to release the inserted DNA from lambda arms) and separated by electrophoresis on a 0.6% Agarose-TAE gel. The large fragments, electroeluted and concentrated as described above, were ligated to SalI-digested and dephosphorylated pUC19 (NEB). The ligation mixture was introduced by transformation into E. coli DH5(alpha) competent cells (BRL) and plated on LB agar containing ampicillin, isopropyl-(Beta)-D-thiogalactoside (IPTG) and 5-Bromo-4-Chloro-3-indolyl-(Beta)-D-galactoside (XGAL). White colonies (with insertions in the (Beta)-galactosidase gene of pUC19) were subjected to standard miniprep procedures to isolate the plasmid, designated pM4,122-3. The full length toxin gene was sequenced by using oligonucleotide

primers made to the "4.5 Kb class" toxin gene and by "walking" with primers made to the sequence of PS81A2.

The plasmid pM4,122-3 contains about 15 Kb of PS81A2 DNA including the 3.522 Kb which encodes the 133,601 dalton endotoxin. The ORF of the PS81A2 toxin gene was isolated from pM4,122-3 and subcloned into the Bacillus shuttle vector pBC1ac as a 5.5 Kb blunt-ended DraIII fragment. E. coli NM522 cells were transformed and plated on LB agar supplemented with ampicillin. The resulting colonies were analyzed by standard miniprep procedures to isolate plasmids that contained the insert. The desired plasmid, pMYC389, contains the coding sequence of the PS81A2 toxin gene.

Example 3 - Cloning of Novel Toxin Gene From Isolate PS81RR1 and Transformation into *Escherichia coli*

Total cellular DNA was prepared from B.t. cells grown to a low optical density ($OD_{600} = 1.0$). The cells were recovered by centrifugation and protoplasted in TES buffer (30 mM Tris-Cl, 10 mM ethylenediaminetetraacetic acid [EDTA], 50 mM NaCl, pH = 8.0) containing 20% sucrose and 50 mg/ml lysozyme. The protoplasts were lysed by addition of sodium dodecyl sulfate (SDS) to a final concentration of 4%. The cellular material was precipitated overnight at 4°C in 100 mM (final concentration) neutral potassium chloride. The supernate was extracted twice with phenol/chloroform (1:1). The DNA was precipitated with ethanol and purified by isopycnic banding on a cesium chloride gradient.

Total cellular DNA from PS81RR1 and B.t.k. HD-1 was digested with EcoRI and separated by electrophoresis on a 0.8% Agarose-TAE-buffered gel. A Southern blot of the gel was probed with the NsiI to NsiI fragment of the toxin gene contained in plasmid pM3,130-7 of NRRL B-18332 and the NsiI to KpnI fragment of the "4.5 Kb class" toxin gene (Kronstad and Whitely [1986])

Gene USA 43:29-40). These two fragments were combined and used as the probe. Results show that hybridizing fragments of PS81RR1 are distinct from those of HD-1. Specifically, a 2.3 Kb hybridizing band in PS81RR1 was detected instead of the 3.8 Kb and 1.8 Kb hybridizing bands seen in HD-1.

5 Two hundred micrograms of PS81RR1 total cellular DNA was digested with EcoRI and separated by electrophoresis on a preparative 0.8% Agarose-TAE gel. The 2.2 Kb to 2.4 Kb region of the gel was cut out and the DNA from it was electroeluted and concentrated using an ELUTIPTM-d (Schleicher and Schuell, Keene, NH) ion exchange column. The isolated EcoRI fragments were
10 ligated to LAMBDA ZAPTM EcoRI arms (Stratagene Cloning Systems, La Jolla, CA) and packaged using Gigapak GOLDTM (Stratagene) extracts. The packaged recombinant phage were plated with E. coli strain BB4 (Stratagene) to give high plaque density. The plaques were screened by standard nucleic acid hybridization procedure with radiolabeled probe. The plaques that hybridized were purified
15 and re-screened at a lower plaque density. The resulting purified phage were grown with R408 M13 helper phage (Stratagene) and the recombinant BlueScriptTM (Stratagene) plasmid was automatically excised and packaged. The "phagemid" was re-infected in XL1-Blue E. coli cells (Stratagene) as part of the automatic excision process. The infected XL1-Blue cells were screened for
20 ampicillin resistance and the resulting colonies were analyzed by standard miniprep procedure to find the desired plasmid. The plasmid, designated pM3,31-3, contains an approximate 2.3 Kb EcoRI insert and was sequenced using Stratagene's T7 and T3 primers plus a set of existing B.t. endotoxin oligonucleotide primers. About 600 bp of the toxin gene was sequenced, and
25 data analysis comparing PS81RR1 to other cloned B.t. endotoxin genes showed that the PS81RR1 sequence was unique. A synthetic oligonucleotide (CGTGGATATGGTGAATCTTATG) was constructed to one of the regions in

the PS81RR1 sequence that was least homologous relative to other existing B.t. endotoxin genes.

PS81RR1 total cellular DNA partially digested with Sau3A and fractionated by electrophoresis into a mixture of 9-23 Kb fragments on a 0.6% agarose-TAE gel was ligated into Lambda GEM™-11 (PROMEGA). The packaged phage at a high titer were plated on P2392 E. coli cells (Stratagene) and screened using the radiolabeled synthetic oligonucleotide (aforementioned) as a nucleic acid hybridization probe. Hybridizing plaques were rescreened at a lower plaque density. A single purified hybridizing plaque was used to infect P2392 E. coli cells in liquid culture for preparation of phage for DNA isolation. DNA was isolated by standard procedures. Preparative amounts of recombinant phage DNA were digested with SalI, to release the inserted DNA from lambda arms, and separated by electrophoresis on a 0.6% Agarose-TAE gel. The large fragments, electroeluted and concentrated as described above, were ligated to SalI-digested and dephosphorylated pUC19 (NEB). The ligation mixture was introduced by transformation into E. coli DH5(alpha) competent cells (BRL) and plated on LB agar containing ampicillin, isopropyl-(Beta)-D-thiogalactoside (IPTG) and 5-Bromo-4-Chloro-3-indolyl-(Beta)-D-galactoside (XGAL). White colonies (with insertions in the (Beta)-galactosidase gene of pUC19) were subjected to standard miniprep procedures to isolate the plasmid, designated pM1,RR1-A. The full length toxin gene was sequenced by using oligonucleotide primers made to the "4.5 Kb class" toxin gene and by "walking" with primers made to the sequence of PS81RR1.

The plasmid pM1,RR1-A contains about 13 Kb of PS81RR1 DNA including the 3.540 Kb which encodes the 133,367 dalton endotoxin. The ORF of the PS81RR1 toxin gene was isolated from pM1,RR1-A on a 3.8 Kb NdeI fragment and ligated into the Bacillus shuttle vector pBC1ac. E. coli NM522 cells were transformed and the resulting colonies were analyzed by standard

miniprep procedures to isolate plasmids that contained the correct insert. The desired plasmid, pMYC390, contains the coding sequence of the PS81RR1 toxin gene.

5 The above cloning procedures were conducted using standard procedures unless otherwise noted.

10 The various methods employed in the preparation of the plasmids and transformation of host organisms are well known in the art. Also, methods for the use of lambda bacteriophage as a cloning vehicle, i.e., the preparation of lambda DNA, in vitro packaging, and transfection of recombinant DNA, are well known in the art. These procedures are all described in Maniatis, T., Fritsch, E.F., and Sambrook, J. (1982) Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Laboratory, New York. Thus, it is within the skill of those in the genetic engineering art to extract DNA from microbial cells, perform restriction enzyme digestions, electrophorese DNA fragments, tail and anneal plasmid and insert DNA, ligate DNA, transform cells, prepare plasmid DNA, electrophorese proteins, and sequence DNA.

15 The restriction enzymes disclosed herein can be purchased from Bethesda Research Laboratories, Gaithersburg, MD, New England Biolabs, Beverly, MA, or Boehringer-Mannheim, Indianapolis, IN. The enzymes are used according to the instructions provided by the supplier.

20 Plasmid pMYC386 containing the B.t. toxin genes, can be removed from the transformed host microbes by use of standard well-known procedures. For example, E. coli NRRL B-18449 can be subjected to cleared lysate isopycnic density gradient procedures, and the like, to recover pMYC386.

25 Example 4 -- Insertion of Toxin Genes Into Plants

The novel genes coding for the novel insecticidal toxins, as disclosed herein, can be inserted into plant cells using the Ti plasmid from Agrobacter

tumefaciens. Plant cells can then be caused to regenerate into plants (Zambryski, P., Joos, H., Gentello, C., Leemans, J., Van Montague, M. and Schell, J [1983] Cell 32:1033-1043). A particularly useful vector in this regard is pEND4K (Klee, H.J., Yanofsky, M.F. and Nester, E.W. [1985] Bio/Technology 3:637-642). This plasmid can replicate both in plant cells and in bacteria and has multiple cloning sites for passenger genes. The toxin gene, for example, can be inserted into the BamHI site of pEND4K, propagated in E. coli, and transformed into appropriate plant cells.

Example 5 – Cloning of Novel *B. thuringiensis* Genes Into Baculoviruses

The novel genes of the invention can be cloned into baculoviruses such as Autographa californica nuclear polyhedrosis virus (AcNPV). Plasmids can be constructed that contain the AcNPV genome cloned into a commercial cloning vector such as pUC8. The AcNPV genome is modified so that the coding region of the polyhedrin gene is removed and a unique cloning site for a passenger gene is placed directly behind the polyhedrin promoter. Examples of such vectors are pGP-B6874, described by Pennock et al. (Pennock, G.D., Shoemaker, C. and Miller, L.K. [1984] Mol. Cell Biol. 4:399-406), and pAC380, described by Smith et al. (Smith, G.E., Summers, M.D. and Fraser, M.J. [1983] Mol Cell Biol. 3:2156-2165). The gene coding for the novel protein toxin of the invention can be modified with BamHI linkers at appropriate regions both upstream and downstream from the coding region and inserted into the passenger site of one of the AcNPV vectors.

As disclosed previously, the nucleotide sequences encoding the novel B.t. toxin genes are shown in Tables 1 and 4. The deduced amino acid sequences are shown in Tables 2 and 5.

It is well known in the art that the amino acid sequence of a protein is determined by the nucleotide sequence of the DNA. Because of the redundancy

of the genetic code, i.e., more than one coding nucleotide triplet (codon) can be used for most of the amino acids used to make proteins, different nucleotide sequences can code for a particular amino acid. Thus, the genetic code can be depicted as follows:

5	Phenylalanine (Phe)	TTK	Histidine (His)	CAK
	Leucine (Leu)	XTY	Glutamine (Gln)	CAJ
	Isoleucine (Ile)	ATM	Asparagine (Asn)	AAK
	Methionine (Met)	ATG	Lysine (Lys)	AAJ
	Valine (Val)	GTL	Aspartic acid (Asp)	GAK
10	Serine (Ser)	QRS	Glutamic acid (Glu)	GAJ
	Proline (Pro)	CCL	Cysteine (Cys)	TGK
	Threonine (Thr)	ACL	Tryptophan (Trp)	TGG
	Alanine (Ala)	GCL	Arginine (Arg)	WGZ
	Tyrosine (Tyr)	TAK	Glycine (Gly)	GGL
15	Termination signal	TAJ		

Key: Each 3-letter deoxynucleotide triplet corresponds to a trinucleotide of mRNA, having a 5'-end on the left and a 3'-end on the right. All DNA sequences given herein are those of the strand whose sequence correspond to the mRNA sequence, with thymine substituted for uracil. The letters stand for the

20 purine or pyrimidine bases forming the deoxynucleotide sequence.

A = adenine

G = guanine

C = cytosine

T = thymine

25 X = T or C if Y is A or G

X = C if Y is C or T

Y = A, G, C or T if X is C

Y = A or G if X is T

W = C or A if Z is A or G

W = C if Z is C or T

Z = A, G, C or T if W is C

Z = A or G if W is A

5 QR = TC if S is A, G, C or T; alternatively

QR = AG if S is T or C

J = A or G

K = T or C

L = A, T, C or G

10 M = A, C or T

The above shows that the novel amino acid sequences of the B.t. toxins can be prepared by equivalent nucleotide sequences encoding the same amino acid sequence of the protein. Accordingly, the subject invention includes such
15 equivalent nucleotide sequences. In addition it has been shown that proteins of identified structure and function may be constructed by changing the amino acid sequence if such changes do not alter the protein secondary structure (Kaiser, E.T. and Kezdy, F.J. [1984] Science 223:249-255). Thus, the subject invention includes mutants of the amino acid sequence depicted herein which do not alter
20 the protein secondary structure, or if the structure is altered, the biological activity is retained to some degree.

Table 1

10	20	30	40	50	60
1 ATGGAGAATA ATATTGAAAA TCAATGCATA CCTTACAATT GTTTAAATAA TCCTGAAGTA					
61 GAGATATTAG GGATTGAAAG GTCAAATAGT AACGTAGCAG CAGAAATCGG CTTGGGGCTT					
121 AGTCGTCTGC TCGTTTCCCG AATTCCACTA GGGGATTTTA TACTTGGCTT GTTTGATGTA					
181 ATATGGGGGG CTATAGGTCC TTCACAATGG GATATATTTT TAGAGCAAAT TGAGCTATTG					
241 ATCGGCCAAA GAATAGAGGA ATTCGCTAGG AATCAGGCCA TTTCTAGATT ACAAGGGCTA					
310	320	330	340	350	360
301 AGCAATCTTT ACCGAATTTA CACAAATGCT TTTAAAAACT GGGAAAGTAGA TCCTACTAAT					
361 CCAGCATTAA GAGAAGAGAT GCGTATTCAA TTTAATGACA TGAACAGTGC TCTTACAACA					
421 GCTATTCTCT TTTTTCAGT TCAAGGTTAT GAAATTCCTC TTTTATCAGT ATATGTTCAA					
481 GCTGCAAATT TACATTTATC GGTTTTGAGA GATGTTTCAG TGTTTGGACA ACCTTGGGGA					
541 TTTGATGTAG CAACAATCAA TAGTCGTTAT AATGATTTAA CTAGGCTTAT TGCGCAATAT					
610	620	630	640	650	660
601 ACTGATTATG CTGTACGTTG GTATAATACG GGGTTAAATC GTTTACCACG TAATGAAGGG					
661 GTACGAGGAT GGGCAAGATT TAATAGGTTT AGAAGAGAGT TAACAATATC AGTATTAGAT					
721 ATTATTTCTT TTTTCCAAAA TTACGATTCT AGATTATATC CAATTCGAC AATCTATCAA					
781 TTAACGCGGG AAGTATATAC AGATCCGGTA ATTAATATAA CTGATTATAG AGTTACCCCA					
841 AGTTTCGAGA GTATTGAAAA TTCAGCTATT AGAAGTCCCC ATCTTATGGA TTTCTTAAAT					
910	920	930	940	950	960
901 AATATAATTA TTGACACTGA TTTAATTAGA GCGGTTCACT ATTGGGCGGG GCATCGTGTA					
961 ACTTCTCATT TTACCGGTAG TTGCGAAGTG ATAAGCTCCC CTCAATACGG GATAACTGCA					
1021 AACGCAGAAC CGAGTCGAAC TATTGCTCCT AGCAGTTTTC CAGGCTCTAA TCTATTTTAT					
1081 AGAACACTAT CAGACCCCTT CTTCGGAAGA TCCGATAATA TTATGCCAAC ATTAGGAATA					
1141 AATGTAGTGC AGGGGGTAGG ATTCATTCAA CCAAATAATG GTGAAGTTCT ATATAGAAGG					
1210	1220	1230	1240	1250	1260
1201 AGAGGAACAG TAGATTCTCT TGATGAGTTG CCAATTGACG GTGAGAATTC ATTAGTTGGA					
1261 TATAGTCATA GATTAAGTCA CGTTACATTA ACCAGGTCGT TATATAATAC TAATATAACT					
1321 AGCTTGCCAA CATTGTTTTG GACACATCAC AGTGCTACTG ATCGAAATAT AATCTATCCG					
1381 GATGTAATTA CACAAATACC ATTGGTAAAA TCATTCTCCC TTACTTCAGG TACCTCTGTA					
1441 GTCAGAGGCC CAGGATTTAC AGGAGGGGAT ATCATCCGAA CTAACTGTA TGGTAATGTA					
1510	1520	1530	1540	1550	1560
1501 CTAAGTATGA GTCTTAATTT TAGTAATACA TCATTACAGC GGTATCCGCT GAGAGTTCGT					
1561 TATGCTGCTT CTCAACAAT GGTATGAGA GTAAATGTTG GAGGGAGTAC TACTTTTGAT					
1621 CRAGGATTCC CTAGTACTAT GAGTGCAAT GGGTCTTTGA CATCTCAATC ATTTAGATTT					
1681 GCAGAAATTC CTGTAGGCAT TAGTACATCT GGCAGTCAA CTGCTGGAAT AAGTATAAGT					
1741 AATAATCCAG GTAGACAAAC GTTTCACCTA GATAGAATTG AATTTATCCC AGTTGATGCA					
1810	1820	1830	1840	1850	1860
1801 ACATTTGAAG CAGAATATGA TTTAGAAAGA GCACAAAAGG CCGTGAATTC GGTGTTTACT					
1861 TCTTCCAATC AAATCGAGTT AAAACAGAT GTGACGGATT ATCATATTGA TCAAGTATCC					
1921 AATTTAGTAG ATTGTTTATC CGATGAATTT TGTCTGGATG AAAAGCGAGA ATTGTCCGAG					
1981 AAAGTCAAAC ATGCGAAGCG ACTCAGTGAT GAGCGGAATT TACTTCAAGA TCCAAACTTC					
2041 AGAGGGATCA ATAGGCAACC AGACCGTGGC TGGAGAGGAA GTACGGATAT TACCATCCAA					
2110	2120	2130	2140	2150	2160
2101 GGAGGAGATG ACCTATTCAA AGAGAATTAC GTCACACTAC CAGGTACCTT TGATGAGTGC					
2161 TATCCAACGT ATTTGTATCA AAAAATAGAT GAGTCGAAAT TAAAGGCTA TAACCGTTAC					
2221 CAATTAAGAG GGTATATCCA AGATAGTCAA GACTTAGAAA TCTATTTAAT TCGCTACAA					
2281 GCAAAACACG AAACAGTAAA TGTACCAGGT ACGGGTTCCT TATGGCGGCT TTCAGTCGAA					
2341 AGTCCAATTG GAAGGTGTGG AGAACCGAAT CCGGTGTGTC CACACCTTGA ATGGAATCCT					

Table 2

	5	10	15
1	Met	Glu	Asn
16	Asn	Asn	Pro
31	Asn	Val	Ala
46	Ser	Arg	Ile
61	Ile	Trp	Gly
76	Gln	Ile	Glu
91	Asn	Gln	Ala
106	Ile	Tyr	Thr
121	Pro	Ala	Leu
136	Ser	Ala	Leu
151	Glu	Ile	Pro
166	Leu	Ser	Val
181	Phe	Asp	Val
196	Leu	Ile	Gly
211	Gly	Leu	Asn
226	Arg	Phe	Asn
241	Ile	Ile	Ser
256	Pro	Thr	Ile
271	Ile	Asn	Ile
286	Glu	Asn	Ser
301	Asn	Ile	Ile
316	Ala	Gly	His
331	Ile	Ser	Ser
346	Arg	Thr	Ile
361	Arg	Thr	Leu
376	Pro	Thr	Leu
391	Pro	Asn	Asn
406	Ser	Leu	Asp
421	Tyr	Ser	His
436	Asn	Thr	Asn
451	Ser	Ala	Thr
466	Ile	Pro	Leu
481	Val	Arg	Gly
496	Val	Asn	Gly
511	Ser	Leu	Gln
526	Thr	Met	Val
541	Gln	Gly	Phe
556	Gln	Ser	Phe
571	Gly	Ser	Gln
586	Gln	Thr	Phe
601	Thr	Phe	Glu
616	Asn	Ser	Leu
631	Val	Thr	Asp
646	Leu	Ser	Asp
661	Lys	Val	Lys
676	Gln	Asp	Pro
691	Trp	Arg	Gly
706	Phe	Lys	Glu
721	Tyr	Pro	Thr
736	Ala	Tyr	Asn
751	Asp	Leu	Glu
766	Val	Asn	Val
781	Ser	Pro	Ile
796	Leu	Glu	Trp
811	Lys	Cys	Ala

Table 2 (continued)

826 Gly Cys Thr Asp Leu Gln Glu Asp Leu Gly Val Trp Val Val Phe
841 Lys Ile Lys Thr Gln Glu Gly Tyr Ala Arg Leu Gly Asn Leu Glu
856 Phe Ile Glu Glu Lys Pro Leu Ile Gly Glu Ala Leu Ser Arg Val
871 Lys Arg Ala Glu Lys Lys Trp Arg Asp Lys Arg Glu Lys Leu Gln
886 Leu Glu Thr Lys Arg Val Tyr Thr Glu Ala Lys Glu Ala Val Asp
901 Ala Leu Phe Val Asp Ser Gln Tyr Asp Arg Leu Gln Ala Asp Thr
916 Asn Ile Gly Met Ile His Ala Ala Asp Arg Leu Val His Gln Ile
931 His Glu Ala Tyr Leu Pro Glu Leu Pro Phe Ile Pro Gly Ile Asn
946 Val Val Ile Phe Glu Glu Leu Glu Asn Arg Ile Ser Thr Ala Leu
961 Ser Leu Tyr Asp Ala Arg Asn Val Ile Lys Asn Gly Asp Phe Asn
976 Asn Gly Leu Ser Cys Trp Asn Val Lys Gly His Val Asp Val Val
991 Glu Gln Asn Asn His Arg Ser Val Leu Val Val Pro Glu Trp Glu
1006 Ala Glu Val Ser Gln Thr Ile Arg Val Cys Pro Gly Arg Gly Tyr
1021 Ile Leu Arg Val Thr Ala Tyr Lys Glu Gly Tyr Gly Glu Gly Cys
1036 Val Thr Ile His Glu Ile Glu Asn Asn Thr Asp Glu Leu Lys Phe
1051 Lys Asn Cys Glu Glu Glu Glu Val Tyr Pro Thr Asp Thr Gly Thr
1066 Cys Asn Asp Tyr Thr Ala His Gln Gly Thr Ala Gly Ser Thr Asp
1081 Ser Cys Asn Ser Arg Asn Ile Arg Tyr Glu Asp Ala Tyr Glu Met
1096 Asn Thr Thr Ala Ser Val Asn Tyr Lys Pro Thr Tyr Glu Glu Glu
1111 Arg Tyr Thr Asp Val Gln Gly Asp Asn His Cys Glu Tyr Asp Arg
1126 Gly Tyr Val Asn Tyr Arg Pro Val Pro Ala Gly Tyr Val Thr Lys
1141 Glu Leu Glu Tyr Phe Pro Glu Thr Asp Lys Val Trp Ile Glu Ile
1156 Gly Glu Thr Glu Gly Lys Phe Ile Val Asp Asn Val Glu Leu Leu
1171 Leu Met Glu Glu

Table 3

5	10	15	20
Met Glu Asn Asn Ile Glu Asn Gln Cys Ile Pro Tyr Asn Cys Leu Asn Asn Pro Glu Val			
ATG GAG AAT AAT ATT GAA AAT CAA TGC ATA CCT TAC AAT TGT TTA AAT AAT CCT GAA GTA			
25	30	35	40
Glu Ile Leu Gly Ile Glu Arg Ser Asn Ser Asn Val Ala Ala Glu Ile Gly Leu Gly Leu			
GAG ATA TTA GGG ATT GAA AGG TCA AAT AGT AAC GTA GCA GCA GAA ATC GGC TTG GGG CTT			
45	50	55	60
Ser Arg Leu Leu Val Ser Arg Ile Pro Leu Gly Asp Phe Ile Leu Gly Leu Phe Asp Val			
AGT CGT CTG CTC GTT TCC CGA ATT CCA CTA GGG GAT TTT ATA CTT GGC TTG TTT GAT GTA			
65	70	75	80
Ile Trp Gly Ala Ile Gly Pro Ser Gln Trp Asp Ile Phe Leu Glu Gln Ile Glu Leu Leu			
ATA TGG GGG GCT ATA GGT CCT TCA CAA TGG GAT ATA TTT TTA GAG CAA ATT GAG CTA TTG			
85	90	95	100
Ile Gly Gln Arg Ile Glu Glu Phe Ala Arg Asn Gln Ala Ile Ser Arg Leu Gln Gly Leu			
ATC GGC CAA AGA ATA GAG GAA TTC GCT AGG AAT CAG GCA ATT TCT AGA TTA CAA GGG CTA			
105	110	115	120
Ser Asn Leu Tyr Arg Ile Tyr Thr Asn Ala Phe Lys Asn Trp Glu Val Asp Pro Thr Asn			
AGC AAT CTT TAC CGA ATT TAC ACA AAT GCT TTT AAA AAC TGG GAA GTA GAT CCT ACT AAT			
125	130	135	140
Pro Ala Leu Arg Glu Glu Met Arg Ile Gln Phe Asn Asp Met Asn Ser Ala Leu Thr Thr			
CCA GCA TTA AGA GAA GAG ATG CGT ATT CAA TTT AAT GAC ATG AAC AGT GCT CTT ACA ACA			
145	150	155	160
Ala Ile Pro Leu Phe Ser Val Gln Gly Tyr Glu Ile Pro Leu Leu Ser Val Tyr Val Gln			
GCT ATT CCT CTT TTT TCA GTT CAA GGT TAT GAA ATT CCT CTT TTA TCA GTA TAT GTT CAA			
165	170	175	180
Ala Ala Asn Leu His Leu Ser Val Leu Arg Asp Val Ser Val Phe Gly Gln Arg Trp Gly			
GCT GCA AAT TTA CAT TTA TCG GTT TTG AGA GAT GTT TCA GTG TTT GGA CAA CGT TGG GGA			
185	190	195	200
Phe Asp Val Ala Thr Ile Asn Ser Arg Tyr Asn Asp Leu Thr Arg Leu Ile Gly Glu Tyr			
TTT GAT GTA GCA ACA ATC AAT AGT CGT TAT AAT GAT TTA ACT AGG CTT ATT GGC GAA TAT			
205	210	215	220
Thr Asp Tyr Ala Val Arg Trp Tyr Asn Thr Gly Leu Asn Arg Leu Pro Arg Asn Glu Gly			
ACT GAT TAT GCT GTA CGT TGG TAT AAT ACG GGG TTA AAT CGT TTA CCA CGT AAT GAA GGG			
225	230	235	240
Val Arg Gly Trp Ala Arg Phe Asn Arg Phe Arg Arg Glu Leu Thr Ile Ser Val Leu Asp			
GTA CGA GGA TGG GCA AGA TTT AAT AGG TTT AGA AGA GAG TTA ACA ATA TCA GTA TTA GAT			
245	250	255	260
Ile Ile Ser Phe Phe Gln Asn Tyr Asp Ser Arg Leu Tyr Pro Ile Pro Thr Ile Tyr Gln			
ATT ATT TCT TTT TTC CAA AAT TAC GAT TCT AGA TTA TAT CCA ATT CCG ACA ATC TAT CAA			
265	270	275	280
Leu Thr Arg Glu Val Tyr Thr Asp Pro Val Ile Asn Ile Thr Asp Tyr Arg Val Thr Pro			
TTA ACG CCG GAA GTA TAT ACA GAT CCG GTA ATT AAT ATA ACT GAT TAT AGA GTT ACC CCA			

Table 3 (continued)

285	290	295	300
Ser Phe Glu Ser Ile Glu Asn Ser Ala Ile Arg Ser Pro His Leu Met Asp Phe Leu Asn			
AGT TTC GAG AGT ATT GAA AAT TCA GGT ATT AGA AGT CCC CAT CTT ATG CAT TTC TTA AAT			
305	310	315	320
Asn Ile Ile Ile Asp Thr Asp Leu Ile Arg Gly Val His Tyr Trp Ala Gly His Arg Val			
AAT ATA ATT ATT GAC ACT GAT TTA ATT AGA GGC GTT CAC TAT TGG GCG GCG CAT CGT GTA			
325	330	335	340
Thr Ser His Phe Thr Gly Ser Ser Gln Val Ile Ser Ser Pro Gln Tyr Gly Ile Thr Ala			
ACT TCT CAT TTT ACC GGT AGT TCG CAA GTG ATA AGC TCC CCG CAA TAC GCG ATA ACT GCA			
345	350	355	360
Asn Ala Glu Pro Ser Arg Thr Ile Ala Pro Ser Thr Phe Pro Gly Leu Asn Leu Phe Tyr			
AAC GCA GAA CCG AGT CGA ACT ATT GGT CCG AGC ACT TTT CCA GGT GTT AAT CTA TTT TAT			
365	370	375	380
Arg Thr Leu Ser Asp Pro Phe Phe Arg Arg Ser Asp Asn Ile Met Pro Thr Leu Gly Ile			
AGA ACA CTA TCA GAC CCG TTC TTC CGA AGA TCC CAT AAT ATT ATG CCA ACA TTA GGA ATA			
385	390	395	400
Asn Val Val Gln Gly Val Gly Phe Ile Gln Pro Asn Asn Gly Glu Val Leu Tyr Arg Arg			
AAT GTA GTG CAG GCG GTA GGA TTC ATT CAA CCA AAT AAT GGT GAA GTT CTA TAT AGA AGG			
405	410	415	420
Arg Gly Thr Val Asp Ser Leu Asp Glu Leu Pro Ile Asp Gly Glu Asn Ser Leu Val Gly			
AGA GGA ACA GTA GAT TCT CTT GAT GAG TTG CCA ATT GAC GGT GAG AAT TCA TTA GTT GGA			
425	430	435	440
Tyr Ser His Arg Leu Ser His Val Thr Leu Thr Arg Ser Leu Tyr Asn Thr Asn Ile Thr			
TAT AGT CAT AGA TTA AGT CAC GTT ACA TTA ACC AGG TCG TTA TAT AAT ACT AAT ATA ACT			
445	450	455	460
Ser Leu Pro Thr Phe Val Trp Thr His His Ser Ala Thr Asp Arg Asn Ile Ile Tyr Pro			
AGC TTG CCA ACA TTT GTT TGG ACA CAT CAC AGT GGT ACT GAT CCA AAT ATA ATC TAT CCG			
465	470	475	480
Asp Val Ile Thr Gln Ile Pro Leu Val Lys Ser Phe Ser Leu Thr Ser Gly Thr Ser Val			
GAT GTA ATT ACA CAA ATA CCA TTG GTA AAA TCA TTC TCC GTT ACT TCA GGT ACC TCT GTA			
485	490	495	500
Val Arg Gly Pro Gly Phe Thr Gly Gly Asp Ile Ile Arg Thr Asn Val Asn Gly Asn Val			
GTG AGA GGC CCA GGA TTT ACA GGA GCG GAT ATC ATC CCA ACT AAC GTT AAT GGT AAT GTA			
505	510	515	520
Leu Ser Met Ser Leu Asn Phe Ser Asn Thr Ser Leu Gln Arg Tyr Arg Val Arg Val Arg			
CTA AGT ATG AGT GTT AAT TTT AGT AAT ACA TCA TTA CAG CCG TAT CCG GTG AGA GTT CGT			
525	530	535	540
Tyr Ala Ala Ser Gln Thr Met Val Met Arg Val Asn Val Gly Gly Ser Thr Thr Phe Asp			
TAT GGT GCT TCT CAA ACA ATG GTC ATG AGA GTA AAT GTT GGA GCG AGT ACT ACT TTT CAT			
545	550	555	560
Gln Gly Phe Pro Ser Thr Met Ser Ala Asn Gly Ser Leu Thr Ser Gln Ser Phe Arg Phe			
CAA GGA TTC CCG AGT ACT ATG AGT GCA AAT GCG TCT TTG ACA TCT CAA TCA TTT AGA TTT			
565	570	575	580
Ala Glu Phe Pro Val Gly Ile Ser Thr Ser Gly Ser Gln Thr Ala Gly Ile Ser Ile Ser			
GCA GAA TTT CCG GTA GCG ATT AGT ACA TCT GCG AGT CAA ACT GGT GGA ATA AGT ATA AGT			

Table 3 (continued)

585	590	595	600
Asn Asn Pro Gly Arg Gln Thr Phe His Leu Asp Arg Ile Glu Phe Ile Pro Val Asp Ala			
AAT AAT CCA GGT AGA CAA ACG TTT CAC TTA GAT AGA ATT GAA TTT ATC CCA GTT GAT GCA			
605	610	615	620
Thr Phe Glu Ala Glu Tyr Asp Leu Glu Arg Ala Gln Lys Ala Val Asn Ser Leu Phe Thr			
ACA TTT GAA GCA GAA TAT GAT TTA GAA AGA GCA CAA AAG GCG GTG AAT TCG CTG TTT ACT			
625	630	635	640
Ser Ser Asn Gln Ile Glu Leu Lys Thr Asp Val Thr Asp Tyr His Ile Asp Gln Val Ser			
TCT TCC AAT CAA ATC GAG TTA AAA ACA GAT GTG ACG GAT TAT CAT ATT GAT CAA GTA TCC			
645	650	655	660
Asn Leu Val Asp Cys Leu Ser Asp Glu Phe Cys Leu Asp Glu Lys Arg Glu Leu Ser Glu			
AAT TTA GTA GAT TGT TTA TCC GAT GAA TTT TGT CTG GAT GAA AAG CGA GAA TTG TCC GAG			
665	670	675	680
Lys Val Lys His Ala Lys Arg Leu Ser Asp Glu Arg Asn Leu Leu Gln Asp Pro Asn Phe			
AAA GTC AAA CAT GCG AAG CGA CTC AGT GAT GAG CGG AAT TTA CTT CAA GAT CCA AAC TTC			
685	690	695	700
Arg Gly Ile Asn Arg Gln Pro Asp Arg Gly Trp Arg Gly Ser Thr Asp Ile Thr Ile Gln			
AGA GGG ATC AAT AGG CAA CCA GAC CGT GGC TGG AGA GGA AGT ACG GAT ATT ACC ATC CAA			
705	710	715	720
Gly Gly Asp Asp Val Phe Lys Glu Asn Tyr Val Thr Leu Pro Gly Thr Phe Asp Glu Cys			
GGA GGA GAT GAC GTA TTC AAA GAG AAT TAC GTC ACA CTA CCA GGT ACC TTT GAT GAG TGC			
725	730	735	740
Tyr Pro Thr Tyr Leu Tyr Gln Lys Ile Asp Glu Ser Lys Leu Lys Ala Tyr Asn Arg Tyr			
TAT CCA ACG TAT TTG TAT CAA AAA ATA GAT GAG TCG AAA TTA AAA GCC TAT AAC CGT TAC			
745	750	755	760
Gln Leu Arg Gly Tyr Ile Glu Asp Ser Gln Asp Leu Glu Ile Tyr Leu Ile Arg Tyr Asn			
CAA TTA AGA GGG TAT ATC GAA GAT AGT CAA GAC TTA GAA ATC TAT TTA ATT CGC TAC AAT			
765	770	775	780
Ala Lys His Glu Thr Val Asn Val Pro Gly Thr Gly Ser Leu Trp Pro Leu Ser Val Glu			
GCA AAA CAC GAA ACA GTA AAT GTA CCA GGT ACG GGT TCC TTA TGG CCG CTT TCA GTC GAA			
785	790	795	800
Ser Pro Ile Gly Arg Cys Gly Glu Pro Asn Arg Cys Val Pro His Leu Glu Trp Asn Pro			
AGT CCA ATT GCA ACG TGT GGA GAA CCG AAT CCG TGT GTG CCA CAC CTT GAA TGG AAT CCT			
805	810	815	820
Asp Leu Asp Cys Ser Cys Arg Asp Gly Glu Lys Cys Ala His His Ser His His Phe Ser			
GAT TTA GAT TGT TCC TGC AGA GAC GGG GAA AAA TGT GCA CAT CAT TCC CAT CAT TTC TCC			
825	830	835	840
Leu Asp Ile Asp Val Gly Cys Thr Asp Leu Gln Glu Asp Leu Gly Val Trp Val Val Phe			
TTG GAC ATT GAT GTT GGA TCC ACA GAC TTG CAA GAG GAT CTA GGC GTG TGG GTT GTA TTC			
845	850	855	860
Lys Ile Lys Thr Gln Glu Gly Tyr Ala Arg Leu Gly Asn Leu Glu Phe Ile Glu Glu Lys			
AAG ATT AAG ACG CAG GAA GGT TAT GCA AGA TTA GGA AAT CTG GAA TTT ATC GAA GAG AAA			
865	870	875	880
Pro Leu Ile Gly Glu Ala Leu Ser Arg Val Lys Arg Ala Glu Lys Lys Trp Arg Asp Lys			
CCA TTA ATT GGA GAA GCA CTG TGT CGT GTG AAG AGA GCG GAA AAA AAA TCG AGA GAC AAA			

Table 4

10	20	30	40	50	60
1	ATGGAGATAA	TGAATAATCA	GAATCAATGC	GTTCCTTATA	ACTGTTTGAA
61	ATTGAAATAT	TAGAAGGAGA	AAGAATAGAA	ACTGGTTACA	CCCCAATAGA
121	TCGCTAACGC	AATTTCTGTT	GAGTGAATTT	GTCCAGGTG	CTGGTTTGT
181	ATTGATTTAA	TATGGGGGTT	TGTGGGTCCC	TCTCAATGGG	ATGCATTCT
241	GAACAGTTAA	TTAACCAAAG	AATAGAGGAA	TTCCGTAGGA	ACCAAGCAAT
					TTCTAGATTA
310	320	330	340	350	360
301	GAAGGGCTAA	GCAACCTTTA	TCAAATTTAC	GCAGAAGCTT	TTAGAGAGTG
361	CCTACTAATC	CAGCATTAA	AGAAGAGATG	CGTATTCAGT	TCAATGACAT
421	CTTACAACCG	CTATTCTCT	TTTTACAGTT	CAAAATTATC	AAGTACCTCT
481	TATGTTCAAG	CTGCAAAATT	ACATTTATCG	GTTTTGAGAG	ATGTTTCAGT
541	CGTTGGGGAT	TTGATGTAGC	AACAATCAAT	AGTCGTTATA	ATGATTTAAC
					TAGGCTTATT
610	620	630	640	650	660
601	GGCACCTATA	CAGATTATGC	TGTACGCTGG	TATAATACGG	GATTAGAACG
661	CCGGATTCTA	CAGATTGGGT	AAGGTATAAT	CAATTTAGAA	GAGAGCTAAC
721	TTAGATATCG	TTTCTCTGTT	CCCGAACTAT	GATAGTAGAA	CGTATCCAAT
781	TCCCAATTA	CTAGAGAAAT	TTATACAAAC	CCAGTATTAG	AAAAATTTGA
841	CGTGGAAATG	CTCAGAGAAT	AGAACAGAAT	ATTAGGCAAC	CACATCTTAT
					GGATCTCCTT
910	920	930	940	950	960
901	AATAGTATAA	CCATTTATAC	TGATGTGCAT	AGAGGCTTTA	ATTATTGGTC
961	ATAACAGCTT	CTCCTGTCCG	TTTTGGGGGG	CCAGAATTTA	CTTTTCCTAG
1021	ATGGGAAATG	CTGCTCCACC	CGTACTGATC	TCAACTACTG	GTTTGGGGAT
1081	TTATCTTCAC	CTCTTTACAG	AAGAATTATA	CTTGGTTTCAG	GGCCAAATAA
1141	TTTGTCTTG	ATGGAACGGG	ATTTTCTTTT	GGCTCCCTAA	CAGCCGATTT
					ACCTTCTACT
1210	1220	1230	1240	1250	1260
1201	ATATACAGAC	AAAGGGGAAC	GGTCGATTCA	CTAGATGTAA	TACCGGCACA
1261	GTCCGAGCAC	GTGCGGGATT	TAGTCATCGA	TTAAGTCATG	TTACAATGCT
1321	GCTGGAGCAG	TTTACACCTT	GAGAGCTCCA	ACGTTTTCTT	GGCGACATCG
1381	TTCTCTAAC	TAATTCCTTC	ATCACAATC	ACACAGATAC	CTTTAACAAA
1441	CTTGCTCTG	GGACCTCTGT	TGTTAAAGGA	CCAGGATTTA	CAGGAGGAGA
					TATTCTTCGA
1510	1520	1530	1540	1550	1560
1501	AGAACTTCAC	CTGGCCAGAT	TTCAACCTTA	AGAGTGACTA	TTACTGCACC
1561	AGATATCGCG	TAAGAATTGG	CTACGTTTCT	ACTACAAATT	TACAATTECA
1621	GACGGAAGAC	CTATTAATCA	GGGGAATTTT	TCAGCAACTA	TGAGTAGTGG
1681	CAGTCGGGAA	GCTTTAGGAC	TGCAGGTTTT	ACTACTCCGT	TTAACTTTTC
1741	AGTATATTTA	CGTTAAGTGC	TCATGTCTTC	AATTCAGGCA	ATGAAGTTTA
					TATAGATCGA
1810	1820	1830	1840	1850	1860
1801	ATTGAATTTG	TTCCGGCAGA	AGTAACATTT	GAGGCGGAAT	ATGATTTAGA
1861	GAGGCGGTGA	ATGCTCTGTT	TACTTCTTCC	AATCAACTAG	GATTAATAAA
1921	GACTATCATA	TTGATCAAST	GTCCAATCTA	GTCCGAATGTT	TATCCGGTGA
1981	GATGAAAAGA	GAGAATTGTC	CGAGAAAGTC	AAACATGCCA	AGCGACTCAG
2041	AATTTACTTC	AAGACCCAAA	CTTCAGAGGC	ATCAATAGAC	AACCAGACCG
					TGGCTGGAGA
2110	2120	2130	2140	2150	2160
2101	GSCAGTACGG	ATATTACCAT	CCAAGGAGGA	GATGACGTAT	TCAAAGAGAA
2161	CTACCGGGTA	CGTTTAATGA	GTGTTATCCT	ACGTATCTGT	ATCAAAAAAT
2221	AAATTAAGAG	CGTATACCGG	TTACCAATTA	AGAGGGTACA	TCGAGGATAG
2281	GAAATCTATT	TAATTCGGTA	CAATACAAA	CACGAAACAG	TAAATGTGCC
2341	TCCTTATGCG	CGCTTTCAGT	CGAAAAATCA	ATTGGAAGT	GCGGAGAACC
					AAATCGATGC

Table 4 (continued)

2401	2420	2430	2440	2450	2460
GCACCACAAC TTGAATGGAA TCCTGATCTA GATTGTTCTT GCAGAGACGG GGAAAAATGT					
2461 GCACATCACT CCCATCATT CTCCTTGGAC ATTGATATTG GATGTACAGA TTAAATGAG					
2521 AACTTAGGTG TATGGGTGAT ATTCAAAATT AAGACGCAAG ATGGTCACGC AAGACTAGGT					
2581 AATCTAGAGT TTCTCAAGA GAAACCATTA GTAGGCGAAT CGTTAGCAGC CGTGAAGAGA					
2641 GCGGAGAAGA AGTGGAGAGA CAAACGAGAG AAATTGCAAG TGGAAACAAA TATCGTTTAT					

2710	2720	2730	2740	2750	2760
2701 AAAGAGGCAA AAGAATCTGT AGATGCTTTA TTTGTGAAC CTCAATATGA TAGATTACAA					
2761 GCGGATACCG ACATCGCGAT GATTCATGCG GCAGATAAAC GCGTTCATCG AATTCGAGAA					
2821 GCATATCTTC CAGAGTTATC TGTAAATCCG GGTGTCAATG CCGGCATTTT TGAAGAATTA					
2881 GAGGGACGTA TTTTCACAGC CTACTCTTTA TATGATGCCA GAAATGTCT TAAAAATGGC					
2941 GATTTCAATA ATGGCTTATC ATGCTGGAAC GTGAAAGGGC ATGTAGATGT AGAAGAACAA					

3010	3020	3030	3040	3050	3060
3001 AACAAACCACC GTTCGGTTCT TGTGTGCGCG GAATGGGAAG CAGAGGTGTC ACAAGAGGTT					
3061 CGTGTCTGTC CAGGTCTGG CTATATCTTA CGTGTACAG CGTACAAAGA GGGATATGGA					
3121 GAAGGTTGCG TAACGATTCA TGAGATCGAA GACAATACAG ACGAACTGAA ATTCAGCAAC					
3181 TGTGTAGAAG AGGAAGTATA TCCAAACAAC ACGGTAACGT GTAATGATTA TACTGCAAAT					
3241 CAAGAAGAAT ACGGGGCTGC GTACACTTCT CGTAATCGTG GATATGGTGA ATCTTATGAA					

3310	3320	3330	3340	3350	3360
3301 AGTAATTCTT CCATACCAGC TGAGTATGCG CCAGTTTATG AGGAAGCATA TATAGATGGA					
3361 AGAAAAGAGA ATCCTTGTGA ATCTAACAGA GGATATGGGG ATTACACGCC ACTACCAGCT					
3421 GGTATGTGA CAAAAGAATT AGAGTACTTC CCAGAAACCG ATAAGGTATG GATTGAGATC					
3481 GGGGAAACCG AAGGAACATT CATCGTGGAT AGCGTGGAA TACTCCTTAT GGAGGAA*					

Segment 1-*

Table 5

	5	10	15
1	Met	Glu	Ile
16	Leu	Asn	Asp
31	Thr	Gly	Tyr
46	Leu	Leu	Ser
61	Ile	Asp	Leu
76	Phe	Leu	Val
91	Phe	Ala	Arg
106	Leu	Tyr	Gln
121	Pro	Thr	Asn
136	Asp	Met	Asn
151	Gln	Asn	Tyr
166	Asn	Leu	His
181	Arg	Trp	Gly
196	Leu	Thr	Arg
211	Tyr	Asn	Thr
226	Trp	Val	Arg
241	Leu	Asp	Ile
256	Pro	Ile	Arg
271	Pro	Val	Leu
286	Arg	Ile	Glu
301	Asn	Ser	Ile
316	Trp	Ser	Gly
331	Pro	Glu	Phe
346	Pro	Pro	Val
361	Leu	Ser	Ser
376	Asn	Asn	Gln
391	Ala	Ser	Leu
406	Gly	Thr	Val
421	Val	Pro	Ala
436	Met	Leu	Ser
451	Thr	Phe	Ser
466	Pro	Ser	Ser
481	Leu	Gly	Ser
496	Gly	Asp	Ile
511	Arg	Val	Thr
526	Ile	Arg	Tyr
541	Asp	Gly	Arg
556	Ser	Gly	Gly
571	Thr	Thr	Pro
586	Ser	Ala	His
601	Ile	Glu	Phe
616	Leu	Glu	Arg
631	Asn	Gln	Leu
646	Gln	Val	Ser
661	Asp	Glu	Lys
676	Leu	Ser	Asp
691	Ile	Asn	Arg
706	Thr	Ile	Gln
721	Leu	Pro	Gly
736	Lys	Ile	Asp
751	Arg	Gly	Tyr
766	Arg	Tyr	Asn
781	Ser	Leu	Trp
796	Glu	Pro	Asn
811	Asp	Cys	Ser

Table 5 (continued)

826 His Phe Ser Leu Asp Ile Asp Ile Gly Cys Thr Asp Leu Asn Glu
841 Asn Leu Gly Val Trp Val Ile Phe Lys Ile Lys Thr Gln Asp Gly
856 His Ala Arg Leu Gly Asn Leu Glu Phe Leu Glu Glu Lys Pro Leu
871 Val Gly Glu Ser Leu Ala Arg Val Lys Arg Ala Glu Lys Lys Trp
886 Arg Asp Lys Arg Glu Lys Leu Gln Val Glu Thr Asn Ile Val Tyr
901 Lys Glu Ala Lys Glu Ser Val Asp Ala Leu Phe Val Asn Ser Gln
916 Tyr Asp Arg Leu Gln Ala Asp Thr Asp Ile Ala Met Ile His Ala
931 Ala Asp Lys Arg Val His Arg Ile Arg Glu Ala Tyr Leu Pro Glu
946 Leu Ser Val Ile Pro Gly Val Asn Ala Gly Ile Phe Glu Glu Leu
961 Glu Gly Arg Ile Phe Thr Ala Tyr Ser Leu Tyr Asp Ala Arg Asn
976 Val Ile Lys Asn Gly Asp Phe Asn Asn Gly Leu Ser Cys Trp Asn
991 Val Lys Gly His Val Asp Val Glu Glu Gln Asn Asn His Arg Ser
1006 Val Leu Val Val Pro Glu Trp Glu Ala Glu Val Ser Gln Glu Val
1021 Arg Val Cys Pro Gly Arg Gly Tyr Ile Leu Arg Val Thr Ala Tyr
1036 Lys Glu Gly Tyr Gly Glu Gly Cys Val Thr Ile His Glu Ile Glu
1051 Asp Asn Thr Asp Glu Leu Lys Phe Ser Asn Cys Val Glu Glu Glu
1066 Val Tyr Pro Asn Asn Thr Val Thr Cys Asn Asp Tyr Thr Ala Asn
1081 Gln Glu Glu Tyr Gly Gly Ala Tyr Thr Ser Arg Asn Arg Gly Tyr
1096 Gly Glu Ser Tyr Glu Ser Asn Ser Ser Ile Pro Ala Glu Tyr Ala
1111 Pro Val Tyr Glu Glu Ala Tyr Ile Asp Gly Arg Lys Glu Asn Pro
1126 Cys Glu Ser Asn Arg Gly Tyr Gly Asp Tyr Thr Pro Leu Pro Ala
1141 Gly Tyr Val Thr Lys Glu Leu Glu Tyr Phe Pro Glu Thr Asp Lys
1156 Val Trp Ile Glu Ile Gly Glu Thr Glu Gly Thr Phe Ile Val Asp
1171 Ser Val Glu Leu Leu Leu Met Glu Glu

Fragment 1-*

Table 6

5	10	15	20
Met Glu Ile Met Asn Asn Gln Asn Gln Cys Val Pro Tyr Asn Cys Leu Asn Asp Pro Thr			
ATG GAG ATA ATG AAT AAT CAG AAT CAA TGC GTT CCT TAT AAC TGT TTG AAT GAT CCG ACA			
25	30	35	40
Ile Glu Ile Leu Glu Gly Glu Arg Ile Glu Thr Gly Tyr Thr Pro Ile Asp Ile Ser Leu			
ATT GAA ATA TTA GAA GGA GAA AGA ATA GAA ACT GGT TAC ACC CCA ATA GAT ATT TCC TTG			
45	50	55	60
Ser Leu Thr Gln Phe Leu Leu Ser Glu Phe Val Pro Gly Ala Gly Phe Val Leu Gly Leu			
TGG CTA ACG CAA TTT CTG TTG AGT GAA TTT GTC CCA GGT GCT GGG TTT GTA TTA GGT TTA			
65	70	75	80
Ile Asp Leu Ile Trp Gly Phe Val Gly Pro Ser Gln Trp Asp Ala Phe Leu Val Gln Ile			
ATT GAT TTA ATA TGG GGG TTT GTG GGT CCC TCT CAA TGG GAT GCA TTT CTT GTG CAA ATT			
85	90	95	100
Glu Gln Leu Ile Asn Gln Arg Ile Glu Glu Phe Ala Arg Asn Gln Ala Ile Ser Arg Leu			
GAA CAG TTA ATT AAC CAA AGA ATA GAG GAA TTC GGT AGG AAC CAA GCA ATT TCT AGA TTA			
105	110	115	120
Glu Gly Leu Ser Asn Leu Tyr Gln Ile Tyr Ala Glu Ala Phe Arg Glu Trp Glu Ala Asp			
GAA GGG CTA AGC AAC CTT TAT CAA ATT TAC GCA GAA GGT TTT AGA GAG TGG GAA GCA GAT			
125	130	135	140
Pro Thr Asn Pro Ala Leu Thr Glu Glu Met Arg Ile Gln Phe Asn Asp Met Asn Ser Ala			
CCT ACT AAT CCA GCA TTA ACA GAA GAG ATG CGT ATT CAG TTC AAT GAC ATG AAC AGT GGT			
145	150	155	160
Leu Thr Thr Ala Ile Pro Leu Phe Thr Val Gln Asn Tyr Gln Val Pro Leu Leu Ser Val			
CTT ACA ACC GGT ATT CCT CTT TTT ACA GTT CAA AAT TAT CAA GTA CCT CTT CTA TCA GTA			
165	170	175	180
Tyr Val Gln Ala Ala Asn Leu His Leu Ser Val Leu Arg Asp Val Ser Val Phe Gly Gln			
TAT GTT CAA GCT GCA AAT TTA CAT TTA TCG GTT TTG AGA GAT GTT TCA GTG TTT GGA CAA			
185	190	195	200
Arg Trp Gly Phe Asp Val Ala Thr Ile Asn Ser Arg Tyr Asn Asp Leu Thr Arg Leu Ile			
CCT TGG GGA TTT GAT GTA GCA ACA ATC AAT AGT CGT TAT AAT GAT TTA ACT AGG CTT ATT			
205	210	215	220
Gly Thr Tyr Thr Asp Tyr Ala Val Arg Trp Tyr Asn Thr Gly Leu Glu Arg Val Trp Gly			
GGC ACC TAT ACA GAT TAT GGT GTA CCG TGG TAT AAT ACG GGA TTA GAA CGT GTA TGG GGA			
225	230	235	240
Pro Asp Ser Arg Asp Trp Val Arg Tyr Asn Gln Phe Arg Arg Glu Leu Thr Leu Thr Val			
CCG GAT TCT AGA GAT TGG GTA AGG TAT AAT CAA TTT AGA AGA GAG CTA ACA CTA ACT GTA			
245	250	255	260
Leu Asp Ile Val Ser Leu Phe Pro Asn Tyr Asp Ser Arg Thr Tyr Pro Ile Arg Thr Val			
TTA GAT ATC GTT TCT CTG TTC CCG AAC TAT GAT AGT AGA ACG TAT CCA ATT CGA ACA GTT			
265	270	275	280
Ser Gln Leu Thr Arg Glu Ile Tyr Thr Asn Pro Val Leu Glu Asn Phe Asp Gly Ser Phe			
TCC CAA TTA ACT AGA GAA ATT TAT ACA AAC CCA GTA TTA GAA AAT TTT GAT GGT AGT TTT			

Table 6 (continued)

285	290	295	300
Arg Gly Met Ala Gln Arg Ile Glu Gln Asn Ile Arg Gln Pro His Leu Met Asp Leu Leu			
CGT GGA ATG GGT CAG AGA ATA GAA CAG AAT ATT AGG CAA CCA CAT CTT ATG GAT CTC CTT			
305	310	315	320
Asn Ser Ile Thr Ile Tyr Thr Asp Val His Arg Gly Phe Asn Tyr Trp Ser Gly His Gln			
AAT AGT ATA ACC ATT TAT ACT GAT GTG CAT AGA GGC TTT AAT TAT TGG TCA GGA CAT CAA			
325	330	335	340
Ile Thr Ala Ser Pro Val Gly Phe Ala Gly Pro Glu Phe Thr Phe Pro Arg Tyr Gly Thr			
ATA ACA GGT TCT CCT GTC GGT TTT GCG GGG CCA GAA TTT ACT TTT CGT AGA TAT GGA ACC			
345	350	355	360
Met Gly Asn Ala Ala Pro Pro Val Leu Ile Ser Thr Thr Gly Leu Gly Ile Phe Arg Thr			
ATG GGA AAT GGT GGT CCA CCG GTA CTG ATC TCA ACT ACT GGT TTG GGG ATT TTT AGA ACA			
365	370	375	380
Leu Ser Ser Pro Leu Tyr Arg Arg Ile Ile Leu Gly Ser Gly Pro Asn Asn Gln Asn Leu			
TTA TCT TCA CCT CTT TAC AGA AGA ATT ATA CTT GGT TCA GGC CCA AAT AAT CAG AAC CTG			
385	390	395	400
Phe Val Leu Asp Gly Thr Glu Phe Ser Phe Ala Ser Leu Thr Ala Asp Leu Pro Ser Thr			
TTT GTC CTT GAT GGA ACG GAA TTT TCT TTT GCG TCC CTA ACA GGC GAT TTA CCT TCT ACT			
405	410	415	420
Ile Tyr Arg Gln Arg Gly Thr Val Asp Ser Leu Asp Val Ile Pro Pro Gln Asp Asn Ser			
ATA TAC AGA CAA AGG GGA ACG GTC GAT TCA CTA GAT GTA ATA CCG CCA CAG GAT AAT AGT			
425	430	435	440
Val Pro Ala Arg Ala Gly Phe Ser His Arg Leu Ser His Val Thr Met Leu Ser Gln Ala			
GTG CCA CCA CGT GCG GGA TTT AGT CAT CGA TTA AGT CAT GTT ACA ATG CTG AGC CAA GCA			
445	450	455	460
Ala Gly Ala Val Tyr Thr Leu Arg Ala Pro Thr Phe Ser Trp Arg His Arg Ser Ala Glu			
GCT GGA CCA GTT TAC ACC TTG AGA GGT CCA ACG TTT TCT TGG CGA CAT CGT AGT GGT GAA			
465	470	475	480
Phe Ser Asn Leu Ile Pro Ser Ser Gln Ile Thr Gln Ile Pro Leu Thr Lys Ser Ile Asn			
TTG TCT AAC CTA ATT CCT TCA TCA CAA ATC ACA CAG ATA CCT TTA ACA AAG TCT ATT AAT			
485	490	495	500
Leu Gly Ser Gly Thr Ser Val Val Lys Gly Pro Gly Phe Thr Gly Gly Asp Ile Leu Arg			
CTT GCG TCT GCG ACC TCT GTT GTT AAA GGA CCA GGA TTT ACA GGA GGA GAT ATT CTT CGA			
505	510	515	520
Arg Thr Ser Pro Gly Gln Ile Ser Thr Leu Arg Val Thr Ile Thr Ala Pro Leu Ser Gln			
AGA ACT TCA CCT GCG CAG ATT TCA ACC TTA AGA GTG ACT ATT ACT CCA CCA TTA TCA CAA			
525	530	535	540
Arg Tyr Arg Val Arg Ile Arg Tyr Ala Ser Thr Thr Asn Leu Gln Phe His Thr Ser Ile			
AGA TAT CCG GTA AGA ATT CCG TAC GCT TCT ACT ACA AAT TTA CAA TTC CAT ACA TCA ATT			
545	550	555	560
Asp Gly Arg Pro Ile Asn Gln Gly Asn Phe Ser Ala Thr Met Ser Ser Gly Gly Asn Leu			
GAC GGA AGA CCT ATT AAT CAG GCG AAT TTT TCA GCA ACT ATG AGT AGT GCG GGT AAT TTA			
565	570	575	580
Gln Ser Gly Ser Arg Thr Ala Gly Phe Thr Thr Pro Phe Asn Phe Ser Asn Gly Ser			
CAG TCC GGA AGC TTT AGG ACT CCA GGT TTT ACT ACT CCG TTT AAC TTT TCA AAT GGA TCA			

Table 6 (continued)

525	590	595	600
Ser Ile Phe Thr Leu Ser Ala His Val Phe Asn Ser Gly Asn Glu Val Tyr Ile Asp Arg			
AGT ATA TTT ACG TTA AGT GCT CAT GTC TTC AAT TCA GGC AAT GAA GTT TAT ATA GAT CGA			
605	610	615	620
Ile Glu Phe Val Pro Ala Glu Val Thr Phe Glu Ala Glu Tyr Asp Leu Glu Arg Ala Gln			
ATT GAA TTT GTT CCG GCA GAA GTA ACA TTT GAG GCG GAA TAT GAT TTA GAA AGA GCG CAA			
625	630	635	640
Glu Ala Val Asn Ala Leu Phe Thr Ser Ser Asn Gln Leu Gly Leu Lys Thr Asn Val Thr			
GAG GCG GTG AAT GGT CTG TTT ACT TCT TCC AAT CAA CTA GGA TTA AAA ACA AAT GTG ACG			
645	650	655	660
Asp Tyr His Ile Asp Gln Val Ser Asn Leu Val Glu Cys Leu Ser Gly Glu Phe Cys Leu			
GAC TAT CAT ATT GAT CAA GTG TCC AAT CTA GTC GAA TGT TTA TCC GGT GAA TTC TGT CTG			
665	670	675	680
Asp Glu Lys Arg Glu Leu Ser Glu Lys Val Lys His Ala Lys Arg Leu Ser Asp Glu Arg			
GAT GAA AAG AGA GAA TTG TCC GAG AAA GTC AAA CAT GCG AAG CGA CTC AGT GAT GAG CCG			
685	690	695	700
Asn Leu Leu Gln Asp Pro Asn Phe Arg Gly Ile Asn Arg Gln Pro Asp Arg Gly Trp Arg			
AAT TTA CTT CAA GAC CCA AAC TTC AGA GGC ATC AAT AGA CAA CCA GAC CGT GCG TGG AGA			
705	710	715	720
Gly Ser Thr Asp Ile Thr Ile Gln Gly Gly Asp Asp Val Phe Lys Glu Asn Tyr Val Thr			
GGC AGT ACG GAT ATT ACC ATC CAA GGA GGA GAT GAC GTA TTC AAA GAG AAT TAC GTC ACA			
725	730	735	740
Leu Pro Gly Thr Phe Asn Glu Cys Tyr Pro Thr Tyr Leu Tyr Gln Lys Ile Asp Glu Ser			
CTA CCG GGT ACC TTT AAT GAG TGT TAT CCT ACG TAT CTG TAT CAA AAA ATA GAT GAG TCG			
745	750	755	760
Lys Leu Lys Ala Tyr Thr Arg Tyr Gln Leu Arg Gly Tyr Ile Glu Asp Ser Gln Asp Leu			
AAA TTA AAA GCG TAT ACC CGT TAC CAA TTA AGA GGG TAC ATC GAG GAT AGT CAA GAC TTA			
765	770	775	780
Glu Ile Tyr Leu Ile Arg Tyr Asn Thr Lys His Glu Thr Val Asn Val Pro Gly Thr Gly			
GAA ATC TAT TTA ATT CCG TAC AAT ACA AAA CAC GAA ACA GTA AAT GTG CCA GGT ACG GGT			
785	790	795	800
Ser Leu Trp Pro Leu Ser Val Glu Asn Pro Ile Gly Lys Cys Gly Glu Pro Asn Arg Cys			
TCC TTA TGG CCG CTT TCA GTC GAA AAT CCA ATT GGA AAG TCC GGA GAA CCA AAT CGA TCC			
805	810	815	820
Ala Pro Gln Leu Glu Trp Asn Pro Asp Leu Asp Cys Ser Cys Arg Asp Gly Glu Lys Cys			
GCA CCA CAA CTT GAA TGG AAT CCG GAT CTA GAT TGT TCC TCC AGA GAC GCG GAA AAA TGT			
825	830	835	840
Ala His His Ser His His Phe Ser Leu Asp Ile Asp Ile Gly Cys Thr Asp Leu Asn Glu			
GCA CAT CAC TCC CAT CAT TTC TCC TTG GAC ATT GAT ATT GGA TGT ACA GAT TTA AAT GAC			
845	850	855	860
Asn Leu Gly Val Trp Val Ile Phe Lys Ile Lys Thr Gln Asp Gly His Ala Arg Leu Gly			
AAC TTA GGT GTA TGG GTG ATA TTC AAA ATT AAG ACG CAA GAT GGT CAC GCA AGA CTA GGT			
865	870	875	880
Asn Leu Glu Phe Leu Glu Glu Lys Pro Leu Val Gly Glu Ser Leu Ala Arg Val Lys Arg			
AAT CTA GAG TTT CTC GAA GAG AAA CCA TTA GTA GGC GAA TCG TTA CCA CCG GTG AAG AGA			

Table 6 (continued)

885	890	895	900
Ala Glu Lys Lys Trp Arg Asp Lys Arg Glu Lys Leu Gln Val Glu Thr Asn Ile Val Tyr			
GGG GAG AAG AAG TGG AGA GAC AAA CGA GAG AAA TTG CAA GTG GAA ACA AAT ATC GTT TAT			
905	910	915	920
Lys Glu Ala Lys Glu Ser Val Asp Ala Leu Phe Val Asn Ser Gln Tyr Asp Arg Leu Gln			
AAA GAG GCA AAA GAA TCT GTA GAT GCT TTA TTT GTG AAG TCT CAA TAT GAT AGA TTA CAA			
925	930	935	940
Ala Asp Thr Asp Ile Ala Met Ile His Ala Ala Asp Lys Arg Val His Arg Ile Arg Glu			
GGG GAT ACC GAC ATC GCG ATG ATT CAT GCG GCA GAT AAA GCG GTT CAT CGA ATT CGA GAA			
945	950	955	960
Ala Tyr Leu Pro Glu Leu Ser Val Ile Pro Gly Val Asn Ala Gly Ile Phe Glu Glu Leu			
GCA TAT GTT CCA GAG TTA TCT GTA ATT CCG GGT GTC AAT GCG GGC ATT TTT GAA GAA TTA			
965	970	975	980
Glu Gly Arg Ile Phe Thr Ala Tyr Ser Leu Tyr Asp Ala Arg Asn Val Ile Lys Asn Gly			
GAG GGA CGT ATT TTC ACA GCG TAC TCT TTA TAT GAT GCG AGA AAT GTC ATT AAA AAT GCG			
985	990	995	1000
Asp Phe Asn Asn Gly Leu Ser Cys Trp Asn Val Lys Gly His Val Asp Val Glu Glu Gln			
GAT TTC AAT AAT GGC TTA TCA TGC TGG AAC GTG AAA GGG CAT GTA GAT GTA GAA GAA CAA			
1005	1010	1015	1020
Asn Asn His Arg Ser Val Leu Val Val Pro Glu Trp Glu Ala Glu Val Ser Gln Glu Val			
AAC AAC CAC CGT TCG GTT CTT GTT GTC CCG GAA TGG GAA GCA GAG GTG TCA CAA GAG GTT			
1025	1030	1035	1040
Arg Val Cys Pro Gly Arg Gly Tyr Ile Leu Arg Val Thr Ala Tyr Lys Glu Gly Tyr Gly			
CGT GTC TGT CCA GGT CGT GGC TAT ATC CTA CGT GTT ACA GCG TAC AAA GAG GGA TAT GGA			
1045	1050	1055	1060
Glu Gly Cys Val Thr Ile His Glu Ile Glu Asp Asn Thr Asp Glu Leu Lys Phe Ser Asn			
GAA GGT TGC GTA ACC ATT CAT GAG ATC GAA GAC AAT ACA GAC GAA CTG AAA TTC AGC AAC			
1065	1070	1075	1080
Cys Val Glu Glu Glu Val Tyr Pro Asn Asn Thr Val Thr Cys Asn Asp Tyr Thr Ala Asn			
TGT GTA GAA GAG GAA GTA TAT CCA AAC AAC ACG GTA ACG TGT AAT GAT TAT ACT GCA AAT			
1085	1090	1095	1100
Gln Glu Glu Tyr Gly Gly Ala Tyr Thr Ser Arg Asn Arg Gly Tyr Gly Glu Ser Tyr Glu			
CAA GAA GAA TAC GGG GGT GCG TAC ACT TCT CGT AAT CGT GGA TAT GGT GAA TCT TAT GAA			
1105	1110	1115	1120
Ser Asn Ser Ser Ile Pro Ala Glu Tyr Ala Pro Val Tyr Glu Glu Ala Tyr Ile Asp Gly			
AGT AAT TCT TCC ATA CCA GCT GAG TAT GCG CCA GTT TAT GAG GAA CCA TAT ATA GAT GGA			
1125	1130	1135	1140
Arg Lys Glu Asn Pro Cys Glu Ser Asn Arg Gly Tyr Gly Asp Tyr Thr Pro Leu Pro Ala			
AGA AAA GAG AAT CCT TGT GAA TCT AAC AGA GGA TAT GGG GAT TAC ACG CCA CTA CCA GCT			
1145	1150	1155	1160
Gly Tyr Val Thr Lys Glu Leu Glu Tyr Phe Pro Glu Thr Asp Lys Val Trp Ile Glu Ile			
GGT TAT GTG ACA AAA GAA TTA GAG TAC TTC CCA GAA ACC GAT AAG GTA TGG ATT GAG ATC			
1165	1170	1175	
Gly Glu Thr Glu Gly Thr Phe Ile Val Asp Ser Val Glu Leu Leu Leu Met Glu Glu			
GGG GAA ACG GAA GCA ACA TTC ATC GTG GAT ACC GTG GAA TTA CTC CTT ATG GAG GAA			

The claims defining the invention are as follows:

1. A process for controlling lepidopteran insect pests which comprises contacting said insect pests with an insect-controlling effective amount of *Bacillus thuringiensis* PS81A2, having the identifying characteristics of NRRL B-18457, or *Bacillus thuringiensis* PS81RR1, having the identifying characteristics of NRRL B-18458, or mutants thereof which retain the characteristics of the parent strains.
2. The process, according to claim 1, wherein said mutants are asporogenous mutants and/or phage resistant mutants.
3. The process, according to claim 1, wherein said insect pest is contacted with an insect-controlling effective amount of *Bacillus thuringiensis* PS81A2 or PS81RR1, by incorporating said *Bacillus thuringiensis* PS81A2 or PS81RR1 into a bait granule and placing said granule on or in the soil when planting seed of a plant upon which plant insect pest is known to feed.
4. A process for controlling soil-inhabiting insect pests of the order Lepidoptera which comprises
 - (1) preparing a bait granule comprising *Bacillus thuringiensis* PS81A2 or PS81RR1, or mutants thereof which retain the characteristics of the parent strains, or spores or crystals of *Bacillus thuringiensis* PS81A2 or PS81RR1; and
 - (2) placing said bait granule on or in the soil.
5. The process, according to claim 4, wherein said bait granule is applied at the same time corn seed is planted in the soil.
6. The process, according to claims 1 or 4, wherein substantially intact *Bacillus thuringiensis* PS81A2 or PS81RR1 cells, or mutants thereof which retain the characteristics of the parent strains, are treated to prolong the pestidical activity when the substantially intact cells are applied to the environment of a target pest.
7. A composition of matter comprising *Bacillus thuringiensis* PS81A2 or PS81RR1, or mutants thereof which retain the characteristics of the parent strains, or spores or crystals of *Bacillus thuringiensis* PS81A2 or PS81RR1, in association with an insecticide carrier, wherein said mutants are asporogenous mutants and/or phage resistant mutants.
8. The composition of matter according to claim 7, wherein said carrier comprises phagostimulants or attractants.
9. A composition of matter comprising *Bacillus thuringiensis* PS81A2 or PS81RR1, or mutants thereof which retain the characteristics of the parent strains, in association with formulation ingredients applied as a seed coating, wherein said mutants are asporogenous mutants and/or phage resistant mutants.
10. *Bacillus thuringiensis* PS81A2, having the identifying characteristics of NRRL B-18457, or mutants thereof which retain the characteristics of the parent strain, having

activity against insect pests of the order Lepidoptera.

11. *Bacillus thuringiensis* PS81RR1, having the identifying characteristics of NRRL B-18458, or mutants thereof which retain the characteristics of the parent strain, having activity against insect pests of the order Lepidoptera.
- 5 12. Asporogenous and/or phage resistant mutants of *Bacillus thuringiensis* PS81A2 or *Bacillus thuringiensis* PS81RR1 which retain the characteristics of the parent strains.

1 13. DNA encoding a Bacillus thuringiensis toxin having the amino acid
2 sequences shown in Table 2 or Table 5.

1 14. DNA, according to claim 13, having the nucleotide sequences shown
2 in Table 1 or Table 4, respectively.

1 15. Toxin active against lepidopteran insects having the amino acid
2 sequence shown in Table 2 or Table 5, and mutants thereof which do not alter
3 the protein secondary structure, or if the structure is altered, the biological
4 activity is retained to some degree.

1 16. A recombinant DNA transfer vector comprising DNA having all or
2 part of the nucleotide sequence which codes for the amino acid sequence shown
3 in Table 2 or Table 5.

1 17. The DNA transfer vector, according to claim 16, transferred to and
2 replicated in a prokaryotic or eukaryotic host.

1 18. A bacterial host transformed to express a Bacillus thuringiensis toxin
2 having the amino acid sequence shown in Table 2 or Table 5.

1 19. Escherichia coli, according to claim 18, transformed with a plasmid
2 vector containing the Bacillus thuringiensis toxin gene encoding the Bacillus
3 thuringiensis toxin having the amino acid sequence shown in Table 2 or Table 5.

1 20. Escherichia coli (NM522)(pMYC389), having the identifying
2 characteristics of NRRL B-18448, or Escherichia coli (NM522)(pMYC390),

3 having the identifying characteristics of NRRL B-18449, hosts according to claim
4 18.

1 21. A microorganism according to claim 18, which is a species of
2 Pseudomonas, Azotobacter, Erwinia, Serratia, Klebsiella, Rhizobium,
3 Rhodopseudomonas, Methylophilus, Agrobacterium, Acetobacter, Alcaligenes,
4 Bacillus, or Streptomyces.

1 22. A microorganism according to claim 21, wherein said microorganism
2 is pigmented and phylloplane adherent.

1 23. A method for controlling lepidopteran insects which comprises
2 administering to said insects or to the environment of said insects a
3 microorganism according to claim 21.

1 24. A method according to claim 23, wherein said administration is to
2 the rhizosphere.

1 25. A method according to claim ²³24, wherein said administration is to
2 the phylloplane.

1 26. A method according to claim 23, wherein said administration is to a
2 body of water.

1 27. An insecticidal composition comprising insecticide containing
2 substantially intact, treated cells having prolonged pesticidal activity when applied
3 to the environment of a target pest, wherein said insecticide is a polypeptide
4 toxic to lepidopteran insects, is intracellular, and is produced as a result of

5 expression of a transformed microbe capable of expressing the Bacillus
6 thuringiensis toxin having the amino acid sequence shown in Table 2 or Table 5.

1 28. The insecticidal composition, according to claim 27, wherein said
2 treated cells are treated by chemical or physical means to prolong their
3 insecticidal activity in the environment.

1 29. The insecticidal composition, according to claim 28, wherein said
2 cells are prokaryotes or lower eukaryotes.

1 30. The insecticidal composition, according to claim 29, wherein said
2 prokaryotic cells are selected from the group consisting of Enterobacteriaceae,
3 Bacillaceae, Rhizobiaceae, Spirillaceae, Lactobacillaceae, Pseudomonadaceae,
4 Azotobacteraceae, Nitrobacteraceae, and Actinomycetales.

1 31. The insecticidal composition, according to claim 29, wherein said
2 lower eukaryotic cells are selected from the group consisting of Phycomycetes,
3 Ascomycetes, and Basidiomycetes.

1 32. The insecticidal composition, according to claim 27, wherein said cell
2 is a pigmented bacterium, yeast, or fungus.

1 33. Treated, substantially intact unicellular microorganism cells containing
2 an intracellular toxin, which toxin is a result of expression of a Bacillus
3 thuringiensis toxin gene toxic to lepidopteran insects which codes for a
4 polypeptide toxin having the amino acid sequence shown in Table 2 or Table 5,
5 wherein said cells are treated under conditions which prolong the insecticidal
6 activity when said cells are applied to the environment of a target insect.